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- (71) Applicant (*for all designated States except US*): **MALLINCKRODT INC.** [US/US]; 675 McDonnell Boulevard, P.O. Box 5840, St. Louis, MO 63134 (US).
- (72) Inventors; and
- (75) Inventors/Applicants (*for US only*): **ACHILEFU, Samuel** [NG/US]; 3424 San Sevilla, St. Louis, MO 63044 (US). **RAJAGOPALAN, Raghavan** [US/US]; 13031 Vinson Court, Maryland Heights, MO 63043 (US). **DORSHOW, Richard, B.** [US/US]; 11977 Neihaus Lane, St. Louis, MO 63146 (US). **BUGAJ, Joseph, E.** [US/US]; 2916 Kettering Drive, St. Charles, MO 63303 (US).
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(54) Title: NOVEL DYES FOR ORGAN FUNCTION MONITORING

(57) Abstract: Highly hydrophilic indole and benzoindole derivatives that absorb and fluoresce in the visible region of light are disclosed. These compounds are useful for physiological and organ function monitoring. Particularly, the molecules of the invention are useful for optical diagnosis of renal and cardiac diseases and for estimation of blood volume in vivo.

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## **NOVEL DYES FOR ORGAN FUNCTION MONITORING**

### **FIELD OF THE INVENTION**

This invention relates to novel optical probes for use in physiological function monitoring, particularly indole and benzoindole  
5 compounds.

### **BACKGROUND OF THE INVENTION**

Dynamic monitoring of physiological functions of patients at the bedside is highly desirable in order to minimize the risk of acute renal failure brought about by various clinical, physiological, and pathological conditions  
10 (C.A. Rabito, L.S.T. Fang, and A.C. Waltman, Renal function in patients at risk with contrast material-induced acute renal failure: Noninvasive real-time monitoring, *Radiology* 1993, 186, 851-854; N.L. Tilney, and J.M. Lazarus, Acute renal failure in surgical patients: Causes, clinical patterns, and care, *Surgical Clinics of North America*, 1983, 63, 357-377; B.E. VanZe, W.E.  
15 Hoy, and J.R. Jaenike, Renal injury associated with intravenous pyelography in non-diabetic and diabetic patients, *Annals of Internal Medicine*, 1978, 89, 51- 54; S. Lundqvist, G. Edbom, S. Groth, U. Stendahl, and S.-O. Hietala, Iohexol clearance for renal function measurement in gynecologic cancer

patients, *Acta Radiologica*, 1996, 37, 582-586; P. Guesry, L. Kaufman, S. Orlof, J.A. Nelson, S. Swann, and M. Holliday, Measurement of glomerular filtration rate by fluorescent excitation of non-radioactive meglumine iothalamate, *Clinical Nephrology*, 1975, 3, 134-138). This monitoring is

5 particularly important in the case of critically ill or injured patients because a large percentage of these patients face the risk of multiple organ failure (MOF), resulting in death (C.C. Baker et al., Epidemiology of Trauma Deaths,

*American Journal of Surgery*, 1980, 144-150; R.G. Lobenhofer et al.,

Treatment Results of Patients with Multiple Trauma: An Analysis of 3406

10 Cases Treated Between 1972 and 1991 at a German Level I Trauma Center, *Journal of Trauma*, 1995, 38, 70-77). MOF is a sequential failure of lung,

liver, and kidneys, and is incited by one or more severe causes such as acute lung injury (ALI), adult respiratory distress syndrome (ARDS),

hypermetabolism, hypotension, persistent inflammatory focus, or sepsis

15 syndrome. The common histological features of hypotension and shock

leading to MOF include tissue necrosis, vascular congestion, interstitial and cellular edema, hemorrhage, and microthrombi. These changes affect the

lung, liver, kidneys, intestine, adrenal glands, brain, and pancreas, in

descending order of frequency (J. Coalson, *Pathology of Sepsis, Septic*

20 *Shock, and Multiple Organ Failure*. In New Horizons: Multiple Organ Failure,

D.J. Bihari and F.B. Cerra (Eds). *Society of Critical Care Medicine*, Fullerton, CA, 1986, pp. 27-59). The transition from early stages of trauma to clinical

MOF is marked by the extent of liver and renal failure and a change in

mortality risk from about 30% to about 50% (F.B. Cerra, Multiple Organ Failure Syndrome. In New Horizons: Multiple Organ Failure, D.J. Bihari and F.B. Cerra (Eds). *Society of Critical Care Medicine*, Fullerton, CA, 1989, pp. 1-24).

- 5                   Serum creatinine measured at frequent intervals by clinical laboratories is currently the most common way of assessing renal function and following the dynamic changes in renal function which occur in critically ill patients (P.D. Dollan, E.L. Alpen, and G.B. Theil, A clinical appraisal of the plasma concentration and endogenous clearance of creatinine, *American*
- 10 *Journal of Medicine*, 1962, 32, 65-79; J.B. Henry (Ed). Clinical Diagnosis and Management by Laboratory Methods, 17th Edition, W.B. Saunders, Philadelphia, PA, 1984); C.E. Speicher, The right test: A physician's guide to laboratory medicine, W.B. Saunders, Philadelphia, PA, 1989). These values are frequently misleading, since age, state of hydration, renal perfusion,
- 15 muscle mass, dietary intake, and many other clinical and anthropometric variables affect the value. In addition, a single value returned several hours after sampling is difficult to correlate with other important physiologic events such as blood pressure, cardiac output, state of hydration and other specific clinical events (e.g., hemorrhage, bacteremia, ventilator settings and others).
- 20 An approximation of glomerular filtration rate can be made via a 24-hour urine collection, but this requires 24 hours to collect the sample, several more hours to analyze the sample, and a meticulous bedside collection technique. New or repeat data are equally cumbersome to obtain.

Occasionally, changes in serum creatinine must be further adjusted based on the values for urinary electrolytes, osmolality, and derived calculations such as the "renal failure index" or the "fractional excretion of sodium." These require additional samples of serum collected contemporaneously with urine samples and, after a delay, precise calculations. Frequently, dosing of medication is adjusted for renal function and thus can be equally as inaccurate, equally delayed, and as difficult to reassess as the values upon which they are based. Finally, clinical decisions in the critically ill population are often as important in their timing as they are in their accuracy.

Exogenous markers such as inulin, iohexol,  $^{51}\text{Cr}$ -EDTA, Gd-DTPA, or  $^{99\text{m}}\text{Tc}$ -DTPA have been reported to measure the glomerular filtration rate (GFR) (P.L. Choyke, H.A. Austin, and J.A. Frank, Hydrated clearance of gadolinium-DTPA as a measurement of glomerular filtration rate, *Kidney International*, 1992, 41, 1595-1598; M.F. Twedle, X. Zhang, M. Fernandez, P. Wedeking, A.D. Nunn, and H.W. Strauss, A noninvasive method for monitoring renal status at bedside, *Invest. Radiol.*, 1997, 32, 802-805; N. Lewis, R. Kerr, and C. Van Buren, Comparative evaluation of urographic contrast media, inulin, and  $^{99\text{m}}\text{Tc}$ -DTPA clearance methods for determination of glomerular filtration rate in clinical transplantation, *Transplantation*, 1989, 48, 790-796). Other markers such as  $^{123}\text{I}$  and  $^{125}\text{I}$  labeled o-iodohippurate or  $^{99\text{m}}\text{Tc}$ -MAG<sub>3</sub> are used to assess tubular secretion process (W.N. Tauxe, Tubular Function, in *Nuclear Medicine in Clinical Urology and Nephrology*, W.N. Tauxe and E.V. Dubovsky, Editors, pp. 77-105, Appleton Century

Crofts, East Norwalk, 1985; R. Muller-Suur, and C. Muller-Suur, Glomerular filtration and tubular secretion of  $MAG_3$  in rat kidney, *Journal of Nuclear Medicine*, 1989, 30, 1986-1991). However, these markers have several undesirable properties such as the use of radioactivity or *ex-vivo* handling of blood and urine samples. Thus, in order to assess the status and to follow the progress of renal disease, there is a considerable interest in developing a simple, safe, accurate, and continuous method for determining renal function, preferably by non-radioactive procedures. Other organs and physiological functions that would benefit from real-time monitoring include the heart, the liver, and blood perfusion, especially in organ transplant patients.

Hydrophilic, anionic substances are generally recognized to be excreted by the kidneys (F. Roch-Ramel, K. Besseghir, and H. Murer, Renal excretion and tubular transport of organic anions and cations, *Handbook of Physiology, Section 8, Neurological Physiology, Vol. II*, E.E. Windhager, Editor, pp. 2189-2262, Oxford University Press, New York, 1992; D.L. Nosco, and J.A. Beaty-Nosco, Chemistry of technetium radiopharmaceuticals 1: Chemistry behind the development of technetium-99m compounds to determine kidney function, *Coordination Chemistry Reviews*, 1999, 184, 91-123). It is further recognized that drugs bearing sulfonate residues exhibit improved clearance through the kidneys (J. Baldas, J. Bonnyman, Preparation, HPLC studies and biological behavior of technetium-99m and  $^{99m}TcNO$ -radiopharmaceuticals based on quinoline type ligands, *Nucl. Med. Biol.*, 1999, 19, 491-496; L. Hansen, A. Taylor, L., L.G. Marzilli, Synthesis

of the sulfonate and phosphonate derivatives of mercaptoacetyltriglycine. X-ray crystal structure of  $\text{Na}_2[\text{ReO}(\text{mercaptoacetylglcylglycylaminomethane-sulfonate})] \cdot 3\text{H}_2\text{O}$ , *Met.-Based Drugs*, 1994, 1, 31-39).

Assessment of renal function by continuously monitoring the  
5 blood clearance of exogenous optical markers, viz., fluorescein bioconjugates  
derived from anionic polypeptides, has been developed by us and by others  
(R.B. Dorshow, J.E. Bugaj, B.D. Burleigh, J.R. Duncan, M.A. Johnson, and  
W.B. Jones, Noninvasive fluorescence detection of hepatic and renal  
function, *Journal of Biomedical Optics*, 1998, 3, 340-345; M. Sohtell et al.,  
10 FITC-Inulin as a Kidney Tubule Marker in the Rat, *Acta. Physiol. Scand.*,  
1983, 119, 313-316, each of which is expressly incorporated herein by  
reference). The main drawback of high molecular weight polypeptides is  
that they are immunogenic. In addition, large polymers with narrow  
molecular weight distribution are difficult to prepare, especially in large  
15 quantities. Thus, there is a need in the art to develop low molecular weight  
compounds that absorb and/or emit light that can be used for assessing  
renal, hepatic, cardiac and other organ functions.

#### SUMMARY OF THE INVENTION

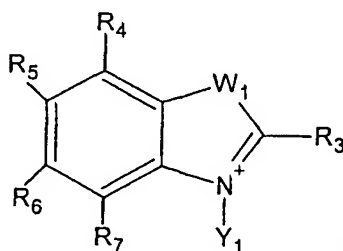
The present invention overcomes these difficulties by  
20 incorporating hydrophilic anionic or polyhydroxy residues in the form of  
sulfates, sulfonates, sulfamates and strategically positioned hydroxyl groups.  
Thus, the present invention is related to novel dyes containing multiple  
hydrophilic moieties and their use as diagnostic agents for assessing organ

function.

The novel compositions of the present invention comprise dyes of Formulas 1 to 6 which are hydrophilic and absorb light in the visible and near infrared regions of the electromagnetic spectrum. The ease of  
5 modifying the clearance pathways of the dyes after *in vivo* administration permits their use for physiological monitoring. Thus, blood protein-binding compounds are useful for angiography and organ perfusion analysis, which is particularly useful in organ transplant and critical ill patients. Predominant  
10 kidney clearance of the dyes enables their use for dynamic renal function monitoring, and rapid liver uptake of the dyes from blood serves as a useful index for the evaluation of hepatic function.

As illustrated in Figures 1-7, these dyes are designed to inhibit aggregation in solution by preventing intramolecular and intermolecular induced hydrophobic interactions.

15 The present invention relates particularly to the novel compounds comprising indoles of the general Formula 1



Formula 1

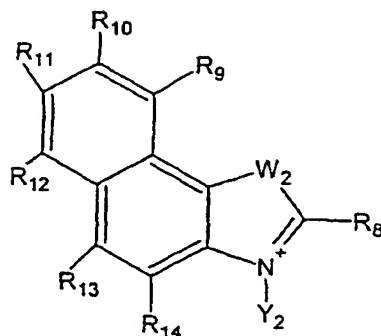


wherein  $R_3$ ,  $R_4$ ,  $R_5$ ,  $R_6$ , and  $R_7$ , and  $Y_1$  are independently selected from the group consisting of -H, C1-C10 alkoxyl, C1-C10 polyalkoxyalkyl, C1-C20 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, saccharides, amino, C1-C10 aminoalkyl, cyano, nitro, halogen, hydrophilic peptides, arylpolysulfonates,

5 C1-C10 alkyl, C1-C10 aryl,  $-\text{SO}_3\text{T}$ ,  $-\text{CO}_2\text{T}$ ,  $-\text{OH}$ ,  $-(\text{CH}_2)_a\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NH}\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{OPO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{OPO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{NHPO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHPO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ , and  $-(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-\text{CH}_2(\text{CH}_2-\text{O}-\text{CH}_2)_c-\text{CH}_2-\text{OH}$ ,  $-(\text{CH}_2)_d-\text{CO}_2\text{T}$ ,  $-\text{CH}_2-(\text{CH}_2-\text{O}-\text{CH}_2)_e-\text{CH}_2-\text{CO}_2\text{T}$ ,  $-(\text{CH}_2)_f-\text{NH}_2$ ,  $-\text{CH}_2-(\text{CH}_2-\text{O}-\text{CH}_2)_g-\text{CH}_2-\text{NH}_2$ ,  $-(\text{CH}_2)_h-\text{N}(\text{R}_a)-(\text{CH}_2)_i-\text{CO}_2\text{T}$ , and  $-(\text{CH}_2)_j-\text{N}(\text{R}_b)-\text{CH}_2-(\text{CH}_2-\text{O}-\text{CH}_2)_k-\text{CH}_2-\text{CO}_2\text{T}$ ;  $W_1$  is selected from the group consisting of  $-\text{CR}_c\text{R}_d$ ,  $-\text{O}-$ ,  $-\text{NR}_c$ ,  $-\text{S}-$ , and  $-\text{Se}-$ ;  $a$ ,  $b$ ,  $d$ ,  $f$ ,  $h$ ,  $i$ , and  $j$  independently vary from 1-10;  $c$ ,  $e$ ,  $g$ , and  $k$  independently vary from 1-100;  $R_a$ ,  $R_b$ ,  $R_c$ , and  $R_d$  are defined in the same manner as  $Y_1$ ;  $T$  is either H or a negative charge.

The present invention also relates to the novel compounds

comprising benzoindoles of general Formula 2

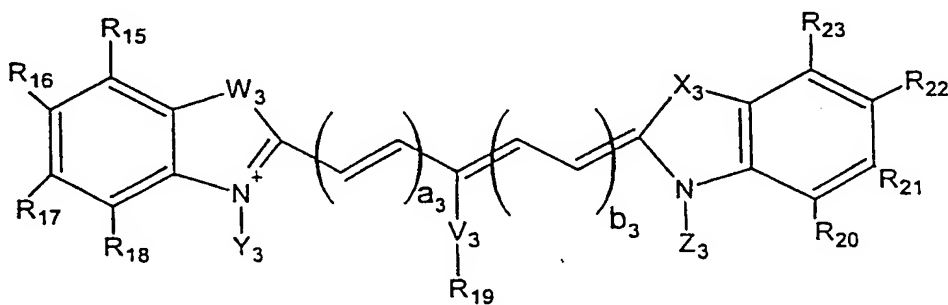


Formula 2

- wherein  $R_8$ ,  $R_9$ ,  $R_{10}$ ,  $R_{11}$ ,  $R_{12}$ ,  $R_{13}$ ,  $R_{14}$ , and  $Y_2$  are independently selected from the group consisting of -H, C1-C10 alkoxyl, C1-C10 polyalkoxyalkyl, C1-C20 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, saccharides, amino, C1-C10 aminoalkyl, cyano, nitro, halogen, hydrophilic peptides, arylpolysulfonates, C1-C10 alkyl, C1-C10 aryl,  $-SO_3T$ ,  $-CO_2T$ ,  $-OH$ ,  $-(CH_2)_aSO_3T$ ,  $-(CH_2)_aOSO_3T$ ,  $-(CH_2)_aNH SO_3T$ ,  $-(CH_2)_aCO_2(CH_2)_bSO_3T$ ,  $-(CH_2)_aOCO(CH_2)_bSO_3T$ ,  $-(CH_2)_aCONH(CH_2)_bSO_3T$ ,  $-(CH_2)_aNHCO(CH_2)_bSO_3T$ ,  $-(CH_2)_aNHCONH(CH_2)_bSO_3T$ ,  $-(CH_2)_aNHCSNH(CH_2)_bSO_3T$ ,  $-(CH_2)_aOCONH(CH_2)_bSO_3T$ ,  $-(CH_2)_aPO_3HT$ ,  $-(CH_2)_aPO_3T_2$ ,  $-(CH_2)_aOPO_3HT$ ,  $-(CH_2)_aOPO_3T_2$ ,  $-(CH_2)_aNHPO_3HT$ ,  $-(CH_2)_aNHPO_3T_2$ ,  $-(CH_2)_aCO_2(CH_2)_bPO_3HT$ ,  $-(CH_2)_aCO_2(CH_2)_bPO_3T_2$ ,  $-(CH_2)_aOCO(CH_2)_bPO_3HT$ ,  $-(CH_2)_aOCO(CH_2)_bPO_3T_2$ , -

- $(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 $(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 $(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 $(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 5  $(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ , and  $-(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-\text{CH}_2(\text{CH}_2\text{-O-CH}_2)_c\text{-CH}_2\text{-OH}$ ,  $-(\text{CH}_2)_d\text{-CO}_2\text{T}$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_e\text{-CH}_2\text{-CO}_2\text{T}$ ,  $-(\text{CH}_2)_f\text{-NH}_2$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_g\text{-CH}_2\text{-NH}_2$ ,  $-(\text{CH}_2)_h\text{-N(R}_a\text{)-(CH}_2)_i\text{-CO}_2\text{T}$ , and  $-(\text{CH}_2)_j\text{-N(R}_b\text{)-CH}_2\text{-(CH}_2\text{-O-CH}_2)_k\text{-CH}_2\text{-CO}_2\text{T}$ ;  $W_2$  is selected from the group consisting of  $-\text{CR}_c\text{R}_d$ ,  $-\text{O-}$ ,  $-\text{NR}_c$ ,  $-\text{S-}$ , and  $-\text{Se-}$ ;  $a$ ,  $b$ ,  $d$ ,  $f$ ,  $h$ ,  $i$ , and  $j$  independently vary from 1-10;  $c$ ,  $e$ ,  $g$ ,  $k$  independently vary from 1-100;  $R_a$ ,  $R_b$ ,  $R_c$ , and  $R_d$  are defined in the same manner as  $Y_2$ ;  $T$  is either H or a negative charge.

The present invention also relates to the novel composition comprising cyanine dyes of general Formula 3



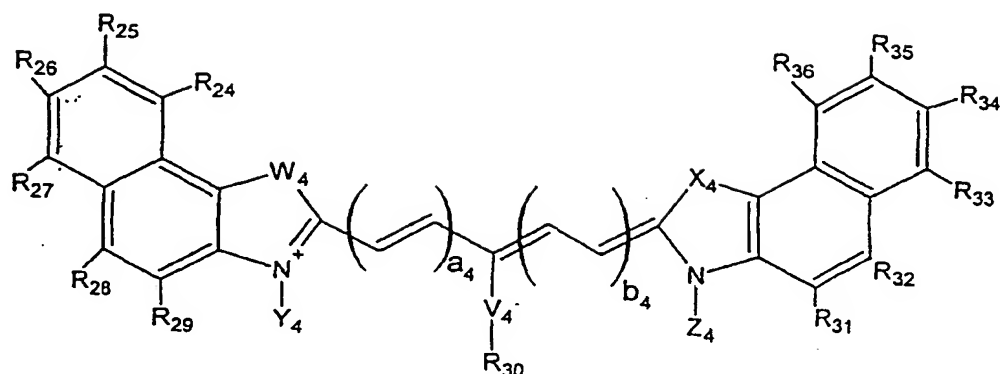
Formula 3

- 15 wherein  $R_{15}$ ,  $R_{16}$ ,  $R_{17}$ ,  $R_{18}$ ,  $R_{19}$ ,  $R_{20}$ ,  $R_{21}$ ,  $R_{22}$ ,  $R_{23}$ ,  $Y_3$ , and  $Z_3$  are independently selected from the group consisting of  $-\text{H}$ , C1-C10 alkoxy, C1-C10

- polyalkoxyalkyl, C1-C20 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, saccharides, amino, C1-C10 aminoalkyl, cyano, nitro, halogen, hydrophilic peptides, arylpolysulfonates, C1-C10 alkyl, C1-C10 aryl,  $-\text{SO}_3\text{T}$ ,  $-\text{CO}_2\text{T}$ ,  $-\text{OH}$ ,  $-(\text{CH}_2)_a\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{SO}_3\text{T}$ , -
- 5  $(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{SO}_3\text{T}$ , -  $(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{SO}_3\text{T}$ , -  $(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{OPO}_3\text{HT}$ , -  $(\text{CH}_2)_a\text{OPO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{NHPO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHPO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{PO}_3\text{HT}$ , -  $(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -
- 10  $(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  $(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  $(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  $(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  $(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ , and  $-(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-\text{CH}_2(\text{CH}_2\text{-O-}$
- 15  $\text{CH}_2)_c\text{-CH}_2\text{-OH}$ ,  $-(\text{CH}_2)_d\text{-CO}_2\text{T}$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_e\text{-CH}_2\text{-CO}_2\text{T}$ ,  $-(\text{CH}_2)_f\text{-NH}_2$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_g\text{-CH}_2\text{-NH}_2$ ,  $-(\text{CH}_2)_h\text{-N(R}_a\text{)-(CH}_2)_i\text{-CO}_2\text{T}$ , and  $-(\text{CH}_2)_j\text{-N(R}_b\text{)-CH}_2\text{-(CH}_2\text{-O-CH}_2)_k\text{-CH}_2\text{-CO}_2\text{T}$ ;  $W_3$  and  $X_3$  are selected from the group consisting of  $-\text{CR}_c\text{R}_d$ ,  $-\text{O-}$ ,  $-\text{NR}_c$ ,  $-\text{S-}$ , and  $-\text{Se-}$ ;  $V_3$  is a single bond or is selected from the group consisting of  $-\text{O-}$ ,  $-\text{S-}$ ,  $-\text{Se-}$ , and  $-\text{NR}_a$ ; a, b, d, f, h, i, and j
- 20 independently vary from 1-10; c, e, g, and k independently vary from 1-100;  $a_3$  and  $b_3$  vary from 0 to 5;  $R_a$ ,  $R_b$ ,  $R_c$ , and  $R_d$  are defined in the same manner as  $Y_3$ ; T is either H or a negative charge.

The present invention further relates to the novel composition

comprising cyanine dyes of general Formula 4

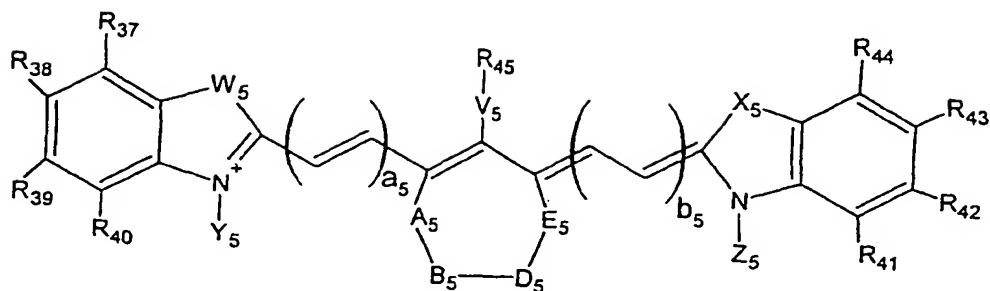


**Formula 4**

- wherein  $R_{24}$ ,  $R_{25}$ ,  $R_{26}$ ,  $R_{27}$ ,  $R_{28}$ ,  $R_{29}$ ,  $R_{30}$ ,  $R_{31}$ ,  $R_{32}$ ,  $R_{33}$ ,  $R_{34}$ ,  $R_{35}$ ,  $R_{36}$ ,  $Y_4$ , and  $Z_4$  are independently selected from the group consisting of -H, C1-C10 alkoxy, C1-C10 polyalkoxyalkyl, C1-C20 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, saccharides, amino, C1-C10 aminoalkyl, cyano, nitro, halogen, hydrophilic peptides, arylpolysulfonates, C1-C10 alkyl, C1-C10 aryl,  $-SO_3T$ ,  $-CO_2T$ ,  $-OH$ ,  $-(CH_2)_aSO_3T$ ,  $-(CH_2)_aOSO_3T$ ,  $-(CH_2)_aNHSO_3T$ ,  $-(CH_2)_aCO_2(CH_2)_bSO_3T$ , -
- 5  $(CH_2)_aOCO(CH_2)_bSO_3T$ ,  $-(CH_2)_aCONH(CH_2)_bSO_3T$ ,  $-(CH_2)_aNHCO(CH_2)_bSO_3T$ , -
- 10  $(CH_2)_aNHCONH(CH_2)_bSO_3T$ ,  $-(CH_2)_aNHCSNH(CH_2)_bSO_3T$ , -

- $(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{OPO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{OPO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{NHPO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHPO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -
- 5  $(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 $(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 $(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 $(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ , and  $-(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-\text{CH}_2(\text{CH}_2\text{-O-CH}_2)_c\text{-CH}_2\text{-OH}$ ,  $-(\text{CH}_2)_d\text{-CO}_2\text{T}$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_e\text{-CH}_2\text{-CO}_2\text{T}$ ,  $-(\text{CH}_2)_f\text{-NH}_2$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_g\text{-CH}_2\text{-NH}_2$ ,  $-(\text{CH}_2)_h\text{-N(R}_a\text{)-(CH}_2)_i\text{-CO}_2\text{T}$ , and  $-(\text{CH}_2)_j\text{-N(R}_b\text{)-CH}_2\text{-(CH}_2\text{-O-CH}_2)_k\text{-CH}_2\text{-CO}_2\text{T}$ ;  $W_4$  and  $X_4$  are selected from the group consisting of  $-\text{CR}_c\text{R}_d$ ,  $-\text{O-}$ ,  $-\text{NR}_c$ ,  $-\text{S-}$ , and  $-\text{Se-}$ ;  $V_4$  is a single bond or is selected from the group consisting of  $-\text{O-}$ ,  $-\text{S-}$ ,  $-\text{Se-}$ , and  $-\text{NR}_a$ ;  $a_4$  and  $b_4$  vary from 0 to 5;  $a$ ,  $b$ ,  $d$ ,  $f$ ,  $h$ ,  $i$ , and  $j$  independently vary from 1-10;  $c$ ,  $e$ ,  $g$ , and  $k$  independently vary from 1-100;  $R_a$ ,  $R_b$ ,  $R_c$ , and  $R_d$  are defined in the same manner as  $Y_4$ ;  $T$  is either H or a negative charge.
- 10  
 15

The present invention also relates to the novel composition comprising cyanine dyes of general Formula 5



Formula 5

wherein  $R_{37}$ ,  $R_{38}$ ,  $R_{39}$ ,  $R_{40}$ ,  $R_{41}$ ,  $R_{42}$ ,  $R_{43}$ ,  $R_{44}$ ,  $R_{45}$ ,  $Y_5$ , and  $Z_5$  are independently selected from the group consisting of -H, C1-C10 alkoxy, C1-C10

- 5 polyalkoxyalkyl, C1-C20 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, saccharides, amino, C1-C10 aminoalkyl, cyano, nitro, halogen, hydrophilic peptides, arylpolysulfonates, C1-C10 alkyl, C1-C10 aryl,  $-\text{SO}_3\text{T}$ ,  $-\text{CO}_2\text{T}$ ,  $-\text{OH}$ ,  $-(\text{CH}_2)_a\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{SO}_3\text{T}$ , -  $(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{SO}_3\text{T}$ , -
- 10  $(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{SO}_3\text{T}$ , -  $(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{OPO}_3\text{HT}$ , -  $(\text{CH}_2)_a\text{OPO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{NHPO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHPO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{PO}_3\text{HT}$ , -  $(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  $(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -
- 15  $(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -

- $-(CH_2)_aCONH(CH_2)_bPO_3T_2$ ,  $-(CH_2)_aNHCO(CH_2)_bPO_3HT$ , -  
 $(CH_2)_aNHCO(CH_2)_bPO_3T_2$ ,  $-(CH_2)_aNHCONH(CH_2)_bPO_3HT$ , -  
 $(CH_2)_aNHCONH(CH_2)_bPO_3T_2$ ,  $-(CH_2)_aNHCSNH(CH_2)_bPO_3HT$ , -  
 $(CH_2)_aNHCSNH(CH_2)_bPO_3T_2$ ,  $-(CH_2)_aOCONH(CH_2)_bPO_3HT$ , and -  
5  $(CH_2)_aOCONH(CH_2)_bPO_3T_2$ ,  $-CH_2(CH_2-O-CH_2)_c-CH_2-OH$ ,  $-(CH_2)_d-CO_2T$ ,  $-CH_2-$   
 $(CH_2-O-CH_2)_e-CH_2-CO_2T$ ,  $-(CH_2)_f-NH_2$ ,  $-CH_2-(CH_2-O-CH_2)_g-CH_2-NH_2$ ,  $-(CH_2)_h-$   
 $N(R_a)-(CH_2)_i-CO_2T$ , and  $-(CH_2)_j-N(R_b)-CH_2-(CH_2-O-CH_2)_k-CH_2-CO_2T$ ;  $W_6$  and  $X_6$   
are selected from the group consisting of  $-CR_cR_d$ ,  $-O-$ ,  $-NR_c$ ,  $-S-$ , and  $-Se$ ;  $V_6$   
is a single bond or is selected from the group consisting of  $-O-$ ,  $-S-$ ,  $-Se-$ , and  
10  $-NR_a$ ;  $D_6$  is a single or a double bond;  $A_6$ ,  $B_6$  and  $E_6$  may be the same or  
different and are selected from the group consisting of  $-O-$ ,  $-S-$ ,  $-Se-$ ,  $-P-$ ,  
 $-NR_a$ ,  $-CR_cR_d$ ,  $CR_c$ , alkyl, and  $-C=O$ ;  $A_6$ ,  $B_6$ ,  $D_6$ , and  $E_6$  may together form a 6  
or 7 membered carbocyclic ring or a 6 or 7 membered heterocyclic ring  
optionally containing one or more oxygen, nitrogen, or sulfur atom;  $a$ ,  $b$ ,  $d$ ,  $f$ ,  
15  $h$ ,  $i$ , and  $j$  independently vary from 1-10;  $c$ ,  $e$ ,  $g$ , and  $k$  independently vary  
from 1-100;  $a_6$  and  $b_6$  vary from 0 to 5;  $R_a$ ,  $R_b$ ,  $R_c$ , and  $R_d$  are defined in the  
same manner as  $Y_6$ ;  $T$  is either H or a negative charge.

The inventive compositions and methods are advantageous  
since they provide a real-time, accurate, repeatable measure of renal  
20 excretion rate using exogenous markers under specific yet changing  
circumstances. This represents a substantial improvement over any currently  
available or widely practiced method, since currently, no reliable, continuous,  
repeatable bedside method for the assessment of specific renal function by



optical methods exists. Moreover, since the inventive method depends solely on the renal elimination of the exogenous chemical entity, the measurement is absolute and requires no subjective interpretation based on age, muscle mass, blood pressure, etc. In fact it represents the nature of renal function

5 in this particular patient, under these particular circumstances, at this precise moment in time.

The inventive compounds and methods provide simple, efficient, and effective monitoring of organ function. The compound is administered and a sensor, either external or internal, is used to detect absorption and/or

10 emission to determine the rate at which the compound is cleared from the blood. By altering the R groups, the compounds may be rendered more organ specific.

#### BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1: Reaction pathway for the preparation of indole

15 derivatives.

Figure 2: Reaction pathway for the preparation of benzoindole derivatives.

Figure 3: Reaction pathway for the preparation of indocarbocyanine derivatives.

20 Figure 4: Reaction pathway for the preparation of benzoindocarbocyanine derivatives.

Figure 5: Reaction pathway for the preparation of robust indocarbocyanine derivatives.

Figure 6: Reaction pathway for the preparation of robust benzoindocarbocyanine derivatives.

Figure 7: Reaction pathway for the preparation of long-wavelength absorbing indocarbocyanine derivatives.

5                    Figure 8a: Absorption spectrum of indoledisulfonate in water.

Figure 8b: Emission spectrum of indoledisulfonate in water.

Figure 9a: Absorption spectrum of indocarbocyaninetetrasulfonate in water.

10                   Figure 9b: Emission spectrum of indocarbocyaninetetrasulfonate in water.

Figure 10a: Absorption spectrum of chloroindocarbocyanine in acetonitrile.

Figure 10b: Emission spectrum of chloroindocarbocyanine in acetonitrile.

15                   Figure 11: Blood clearance profile of carbocyanine-polyaspartic (10 kDa) acid conjugate in a rat.

Figure 12: Blood clearance profile of carbocyanine-polyaspartic (30 kDa) acid conjugate in a rat.

Figure 13: Blood clearance profile of indoledisulfonate in a rat.

20                   Figure 14: Blood clearance profile of carbocyaninetetrasulfonates in a rat.

## DETAILED DESCRIPTION

In one embodiment of the invention, the dyes of the invention

serve as probes for continuous monitoring of renal function, especially for critically ill patients and kidney transplant patients.

In another aspect of the invention, the dyes of the invention are useful for dynamic hepatic function monitoring, especially for critically ill  
 5 patients and liver transplant patients.

In yet another aspect of the invention, the dyes of the invention are useful for real-time determination of cardiac function, especially in patients with cardiac diseases.

In still another aspect of the invention, the dyes of the invention  
 10 are useful for monitoring organ perfusion, especially for critically ill, cancer, and organ transplant patients.

The novel dyes of the present invention are prepared according to the methods well known in the art, as illustrated in general in Figures 1-7 and described for specific compounds in Examples 1-11.

15 In one embodiment, the novel compositions, also called tracers, of the present invention have the Formula 1, wherein  $R_3$ ,  $R_4$ ,  $R_5$ ,  $R_6$  and  $R_7$ , and  $Y_1$  are independently selected from the group consisting of -H, C1-C5 alkoxy, C1-C5 polyalkoxyalkyl, C1-C10 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, mono- and disaccharides, nitro, hydrophilic peptides,  
 20 arylpolysulfonates, C1-C5 alkyl, C1-C10 aryl,  $-\text{SO}_3\text{T}$ ,  $-\text{CO}_2\text{T}$ ,  $-\text{OH}$ ,  $-(\text{CH}_2)_a\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NH}\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-\text{CH}_2(\text{CH}_2\text{-O-CH}_2)_c\text{-CH}_2\text{-OH}$ ,  $-(\text{CH}_2)_d\text{-CO}_2\text{T}$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_e\text{-CH}_2\text{-CO}_2\text{T}$ ,  $-(\text{CH}_2)_f\text{-NH}_2$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_g\text{-CH}_2\text{-NH}_2$ ,  $-(\text{CH}_2)_h\text{-N(R}_3\text{)-}$

$(CH_2)_i-CO_2T$ , and  $-(CH_2)_j-N(R_b)-CH_2-(CH_2-O-CH_2)_k-CH_2-CO_2T$ ;  $W_1$  is selected from the group consisting of  $-CR_cR_d$ ,  $-O-$ ,  $-NR_c$ ,  $-S-$ , and  $-Se$ ;  $a, b, d, f, h, l$ , and  $j$  independently vary from 1-5;  $c, e, g$ , and  $k$  independently vary from 1-20;  $R_a, R_b, R_c$ , and  $R_d$  are defined in the same manner as  $Y_1$ ;  $T$  is a negative charge.

In another embodiment, the novel compositions of the present invention have the general Formula 2, wherein  $R_8, R_9, R_{10}, R_{11}, R_{12}, R_{13}, R_{14}$ , and  $Y_2$  are independently selected from the group consisting of  $-H$ , C1-C5 alkoxyl, C1-C5 polyalkoxyalkyl, C1-C10 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, mono- and disaccharides, nitro, hydrophilic peptides, arylpolysulfonates, C1-C5 alkyl, C1-C10 aryl,  $-SO_3T$ ,  $-CO_2T$ ,  $-OH$ ,  $-(CH_2)_aSO_3T$ ,  $-(CH_2)_aOSO_3T$ ,  $-(CH_2)_aNHSO_3T$ ,  $-(CH_2)_aCO_2(CH_2)_bSO_3T$ ,  $-(CH_2)_aOCO(CH_2)_bSO_3T$ ,  $-CH_2(CH_2-O-CH_2)_c-CH_2-OH$ ,  $-(CH_2)_d-CO_2T$ ,  $-CH_2-(CH_2-O-CH_2)_e-CH_2-CO_2T$ ,  $-(CH_2)_f-NH_2$ ,  $-CH_2-(CH_2-O-CH_2)_g-CH_2-NH_2$ ,  $-(CH_2)_h-N(R_a)-$

$(CH_2)_i-CO_2T$ , and  $-(CH_2)_j-N(R_b)-CH_2-(CH_2-O-CH_2)_k-CH_2-CO_2T$ ;  $W_2$  is selected from the group consisting of  $-CR_cR_d$ ,  $-O-$ ,  $-NR_c$ ,  $-S-$ , and  $-Se$ ;  $a, b, d, f, h, l$ , and  $j$  independently vary from 1-5;  $c, e, g$ , and  $k$  independently vary from 1-20;  $R_a, R_b, R_c$ , and  $R_d$  are defined in the same manner as  $Y_2$ ;  $T$  is a negative charge.

In another embodiment, the novel compositions of the present invention have the general Formula 3, wherein  $R_{15}, R_{16}, R_{17}, R_{18}, R_{19}, R_{20}, R_{21}, R_{22}, R_{23}, Y_3$ , and  $Z_3$  are independently selected from the group consisting of  $-H$ , C1-C5 alkoxyl, C1-C5 polyalkoxyalkyl, C1-C10 polyhydroxyalkyl, C5-C20

polyhydroxyaryl, mono- and disaccharides, nitro, hydrophilic peptides, arylpolysulfonates, C1-C5 alkyl, C1-C10 aryl,  $-\text{SO}_3\text{T}$ ,  $-\text{CO}_2\text{T}$ ,  $-\text{OH}$ ,  $-(\text{CH}_2)_a\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-\text{CH}_2(\text{CH}_2\text{-O-CH}_2)_c\text{-CH}_2\text{-OH}$ ,  $-(\text{CH}_2)_d\text{-CO}_2\text{T}$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_e\text{-CH}_2\text{-CO}_2\text{T}$ ,  $-(\text{CH}_2)_f\text{-NH}_2$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_g\text{-CH}_2\text{-NH}_2$ ,  $-(\text{CH}_2)_h\text{-N(R}_a\text{)-(CH}_2)_i\text{-CO}_2\text{T}$ , and  $-(\text{CH}_2)_j\text{-N(R}_b\text{)-CH}_2\text{-(CH}_2\text{-O-CH}_2)_k\text{-CH}_2\text{-CO}_2\text{T}$ ;  $W_3$  and  $X_3$  are selected from the group consisting of  $-\text{CR}_c\text{R}_d$ ,  $-\text{O-}$ ,  $-\text{NR}_c$ ,  $-\text{S-}$ , and  $-\text{Se-}$ ;  $V_3$  is a single bond or is selected from the group consisting of  $-\text{O-}$ ,  $-\text{S-}$ ,  $-\text{Se-}$ , and  $-\text{NR}_a$ ;  $a$ ,  $b$ ,  $d$ ,  $f$ ,  $h$ ,  $i$ , and  $j$  independently vary from 1-5;  $c$ ,  $e$ ,  $g$ , and  $k$  independently vary from 1-50;  $a_3$  and  $b_3$  vary from 0 to 5;  $R_a$ ,  $R_b$ ,  $R_c$ , and  $R_d$  are defined in the same manner as  $Y_3$ ;  $T$  is either H or a negative charge.

In another embodiment, the novel compositions of the present invention have the general Formula 4, wherein  $R_{24}$ ,  $R_{25}$ ,  $R_{26}$ ,  $R_{27}$ ,  $R_{28}$ ,  $R_{29}$ ,  $R_{30}$ ,  $R_{31}$ ,  $R_{32}$ ,  $R_{33}$ ,  $R_{34}$ ,  $R_{35}$ ,  $R_{36}$ ,  $Y_4$ , and  $Z_4$  are independently selected from the group consisting of  $-\text{H}$ , C1-C5 alkoxyl, C1-C5 polyalkoxyalkyl, C1-C10 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, mono- and disaccharides, nitro, hydrophilic peptides, arylpolysulfonates, C1-C5 alkyl, C1-C10 aryl,  $-\text{SO}_3\text{T}$ ,  $-\text{CO}_2\text{T}$ ,  $-\text{OH}$ ,  $-(\text{CH}_2)_a\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-\text{CH}_2(\text{CH}_2\text{-O-CH}_2)_c\text{-CH}_2\text{-OH}$ ,  $-(\text{CH}_2)_d\text{-CO}_2\text{T}$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_e\text{-CH}_2\text{-CO}_2\text{T}$ ,  $-(\text{CH}_2)_f\text{-NH}_2$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_g\text{-CH}_2\text{-NH}_2$ ,  $-(\text{CH}_2)_h\text{-N(R}_a\text{)-(CH}_2)_i\text{-CO}_2\text{T}$ , and  $-(\text{CH}_2)_j\text{-N(R}_b\text{)-CH}_2\text{-(CH}_2\text{-O-CH}_2)_k\text{-CH}_2\text{-CO}_2\text{T}$ ;  $W_4$  and  $X_4$  are selected from the group consisting of  $-\text{CR}_c\text{R}_d$ ,  $-\text{O-}$ ,  $-\text{NR}_c$ ,  $-\text{S-}$ , and  $-\text{Se-}$ ;  $V_4$  is a single bond or is selected from the group consisting of -

O-, -S-, -Se-, and -NR<sub>a</sub>; a<sub>4</sub> and b<sub>4</sub> vary from 0 to 5; a, b, d, f, h, i, and j independently vary from 1-5; c, e, g, and k independently vary from 1-50; R<sub>a</sub>, R<sub>b</sub>, R<sub>c</sub>, and R<sub>d</sub> are defined in the same manner as Y<sub>4</sub>; T is either H or a negative charge.

- 5 In another embodiment, the novel compositions of the present invention have the general Formula 5, wherein R<sub>37</sub>, R<sub>38</sub>, R<sub>39</sub>, R<sub>40</sub>, R<sub>41</sub>, R<sub>42</sub>, R<sub>43</sub>, R<sub>44</sub>, R<sub>45</sub>, Y<sub>5</sub>, and Z<sub>5</sub> are independently selected from the group consisting of - H, C1-C5 alkoxy, C1-C5 polyalkoxyalkyl, C1-C10 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, mono- and disaccharides, nitro, hydrophilic peptides,
- 10 arylpolysulfonates, C1-C5 alkyl, C1-C10 aryl, -SO<sub>3</sub>T, -CO<sub>2</sub>T, -OH, - (CH<sub>2</sub>)<sub>a</sub>SO<sub>3</sub>T, -(CH<sub>2</sub>)<sub>a</sub>OSO<sub>3</sub>T, -(CH<sub>2</sub>)<sub>a</sub>NHSO<sub>3</sub>T, -(CH<sub>2</sub>)<sub>a</sub>CO<sub>2</sub>(CH<sub>2</sub>)<sub>b</sub>SO<sub>3</sub>T, - (CH<sub>2</sub>)<sub>a</sub>OCO(CH<sub>2</sub>)<sub>b</sub>SO<sub>3</sub>T, -CH<sub>2</sub>(CH<sub>2</sub>-O-CH<sub>2</sub>)<sub>c</sub>-CH<sub>2</sub>-OH, -(CH<sub>2</sub>)<sub>d</sub>-CO<sub>2</sub>T, -CH<sub>2</sub>-(CH<sub>2</sub>-O-CH<sub>2</sub>)<sub>e</sub>-CH<sub>2</sub>-CO<sub>2</sub>T, -(CH<sub>2</sub>)<sub>f</sub>-NH<sub>2</sub>, -CH<sub>2</sub>-(CH<sub>2</sub>-O-CH<sub>2</sub>)<sub>g</sub>-CH<sub>2</sub>-NH<sub>2</sub>, -(CH<sub>2</sub>)<sub>h</sub>-N(R<sub>a</sub>)-(CH<sub>2</sub>)<sub>i</sub>-CO<sub>2</sub>T, and -(CH<sub>2</sub>)<sub>j</sub>-N(R<sub>b</sub>)-CH<sub>2</sub>-(CH<sub>2</sub>-O-CH<sub>2</sub>)<sub>k</sub>-CH<sub>2</sub>-CO<sub>2</sub>T; W<sub>5</sub> and X<sub>5</sub> are
- 15 selected from the group consisting of -CR<sub>c</sub>R<sub>d</sub>, -O-, -NR<sub>c</sub>, -S-, and -Se; V<sub>5</sub> is a single bond or is selected from the group consisting of -O-, -S-, -Se-, and -NR<sub>a</sub> D<sub>5</sub> is a single or a double bond; A<sub>5</sub>, B<sub>5</sub> and E<sub>5</sub> may be the same or different and are selected from the group consisting of -O-, -S-, -NR<sub>a</sub>, -CR<sub>c</sub>R<sub>d</sub>, CR<sub>c</sub>, and alkyl; A<sub>5</sub>, B<sub>5</sub>, D<sub>5</sub>, and E<sub>5</sub> may together form a 6 or 7 membered
- 20 carbocyclic ring or a 6 or 7 membered heterocyclic ring optionally containing one or more oxygen, nitrogen, or sulfur atom; a, b, d, f, h, i, and j independently vary from 1-5; c, e, g, and k independently vary from 1-50; a<sub>5</sub> and b<sub>5</sub> vary from 0 to 5; R<sub>a</sub>, R<sub>b</sub>, R<sub>c</sub>, and R<sub>d</sub> are defined in the same manner as

$Y_5$ ; T is either H or a negative charge.

In yet another embodiment, the novel compositions of the present invention have the general Formula 6, wherein  $R_{46}$ ,  $R_{47}$ ,  $R_{48}$ ,  $R_{49}$ ,  $R_{50}$ ,  $R_{51}$ ,  $R_{52}$ ,  $R_{53}$ ,  $R_{54}$ ,  $R_{55}$ ,  $R_{56}$ ,  $R_{57}$ ,  $R_{58}$ ,  $Y_6$ , and  $Z_6$  are independently selected

5 from the group consisting of -H, C1-C5 alkoxy, C1-C5 polyalkoxyalkyl, C1-C10 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, mono- and disaccharides, nitro, hydrophilic peptides, arylpolysulfonates, C1-C5 alkyl, C1-C10 aryl, - $SO_3T$ , - $CO_2T$ , -OH,  $-(CH_2)_aSO_3T$ ,  $-(CH_2)_aOSO_3T$ ,  $-(CH_2)_aNH_2SO_3T$ , - $(CH_2)_aCO_2(CH_2)_bSO_3T$ ,  $-(CH_2)_aOCO(CH_2)_bSO_3T$ ,  $-CH_2(CH_2-O-CH_2)_c-CH_2-OH$ , - $(CH_2)_d-CO_2T$ ,  $-CH_2-(CH_2-O-CH_2)_e-CH_2-CO_2T$ ,  $-(CH_2)_f-NH_2$ ,  $-CH_2-(CH_2-O-CH_2)_g-$

10  $CH_2-NH_2$ ,  $-(CH_2)_h-N(R_a)-(CH_2)_i-CO_2T$ , and  $-(CH_2)_j-N(R_b)-CH_2-(CH_2-O-CH_2)_k-CH_2-CO_2T$ ;  $W_6$  and  $X_6$  are selected from the group consisting of  $-CR_cR_d$ , -O-, - $NR_c$ , -S-, and -Se;  $V_6$  is a single bond or is selected from the group consisting of -O-, -S-, -Se-, and - $NR_a$ ;  $D_6$  is a single or a double bond;  $A_6$ ,  $B_6$  and  $E_6$  may

15 be the same or different and are selected from the group consisting of -O-, -S-, - $NR_a$ ,  $-CR_cR_d$ ,  $CR_c$ , and alkyl;  $A_6$ ,  $B_6$ ,  $D_6$ , and  $E_6$  may together form a 6 or 7 membered carbocyclic ring or a 6 or 7 membered heterocyclic ring optionally containing one or more oxygen, nitrogen, or sulfur atom; a, b, d, f, h, i, and j independently vary from 1-5; c, e, g, and k independently vary

20 from 1-50;  $a_5$  and  $b_5$  vary from 0 to 5;  $R_a$ ,  $R_b$ ,  $R_c$ , and  $R_d$  are defined in the same manner as  $Y_6$ ; T is either H or a negative charge.

The dosage of the tracers may vary according to the clinical procedure contemplated and generally ranges from 1 picomolar to 100

millimolar. The tracers may be administered to the patient by any suitable method, including intravenous, intraperitoneal, or subcutaneous injection or infusion, oral administration, transdermal absorption through the skin, or by inhalation. The detection of the tracers is achieved by optical fluorescence, absorbance, or light scattering methods known in the art (Muller et al. Eds, Medical Optical Tomography, SPIE Volume IS11, 1993, which is expressly incorporated herein by reference) using invasive or non-invasive probes such as endoscopes, catheters, ear clips, hand bands, surface coils, finger probes, and the like. Physiological function is correlated with the clearance profiles and rates of these agents from body fluids (R.B. Dorshow et al., Non-Invasive Fluorescence Detection of Hepatic and Renal Function, *Bull. Am. Phys. Soc.* 1997, 42, 681, which is expressly incorporated by reference herein).

The organ functions can be assessed either by the differences in the manner in which the normal and impaired cells remove the tracer from the bloodstream, by measuring the rate or accumulation of these tracers in the organs or tissues, or by obtaining tomographic images of the organs or tissues. Blood pool clearance may be measured non-invasively from convenient surface capillaries such as those found in an ear lobe or a finger, for example, using an ear clip or finger clip sensor, or may be measured invasively using an endovascular catheter. Accumulation of the tracer within the cells of interest can be assessed in a similar fashion. The clearance of the tracer dyes may be determined by selecting excitation wavelengths and filters for the emitted photons. The concentration-time curves may be



analyzed in real time by a microprocessor. In order to demonstrate feasibility of the inventive compounds to monitor organ function, a non-invasive absorbance or fluorescence detection system to monitor the signal emanating from the vasculature infused with the compounds is used. Indole derivatives, such as those of Formulas 1-6, fluoresce at a wavelength between 350 nm and 1300 nm when excited at the appropriate wavelength as is known to, or readily determined by, one skilled in the art.

In addition to the noninvasive techniques, a modified pulmonary artery catheter can be used to make the necessary measurements (R.B. Dorshow, J.E. Bugaj, S.A. Achilefu, R. Rajagopalan, and A.H. Combs, Monitoring Physiological Function by Detection of Exogenous Fluorescent Contrast Agents, in *Optical Diagnostics of Biological Fluids IV*, A. Priezzhev and T. Asakura, Editors, Proceedings of SPIE 1999, 3599, 2-8, which is expressly incorporated by reference herein). Currently, pulmonary artery catheters measure only intravascular pressures, cardiac output and other derived measures of blood flow. Critically ill patients are managed using these parameters, but rely on intermittent blood sampling and testing for assessment of renal function. These laboratory parameters represent discontinuous data and are frequently misleading in many patient populations. Yet, importantly, they are relied upon heavily for patient assessment, treatment decisions, and drug dosing.

The modified pulmonary artery catheter incorporates an optical sensor into the tip of a standard pulmonary artery catheter. This wavelength

specific optical sensor can monitor the renal function specific elimination of an optically detectable chemical entity. Thus, by a method analogous to a dye dilution curve, real-time renal function can be monitored by the disappearance of the optically detected compound. Modification of a standard pulmonary artery catheter only requires making the fiber optic sensor wavelength specific, as is known to one skilled in this art. Catheters that incorporate fiber optic technology for measuring mixed venous oxygen saturation currently exist.

- The present invention may be used for rapid bedside evaluation of renal function and also to monitor the efficiency of hemodialysis. The invention is further demonstrated by the following examples. Since many modifications, variations, and changes in detail may be made to the described embodiments, it is intended that all matter in the foregoing description and shown in the accompanying drawings be interpreted as illustrative and not in a limiting sense.

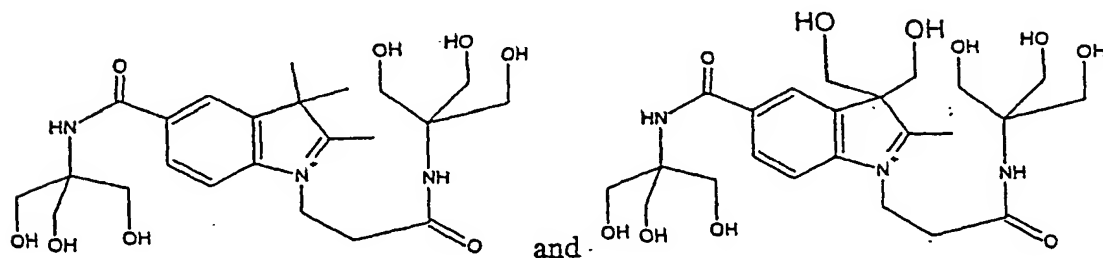
### EXAMPLE 1

#### Synthesis of indole disulfonate (Figure 1, Compound 5, $Y_7 = SO_3^-$ ; $X_7 = H$ ; $n = 1$ )

A mixture of 3-methyl-2-butanone (25.2 mL), and p-hydrazinobenzenesulfonic acid (15 g) in acetic acid (45 mL) was heated at 110°C for 3 hours. After reaction, the mixture was allowed to cool to room temperature and ethyl acetate (100 mL) was added to precipitate the product, which was filtered and washed with ethyl acetate (100 mL). The

intermediate compound, 2,3,3-trimethylindolenium-5-sulfonate (Figure 1, compound 3) was obtained as a pink powder in 80% yield. A portion of compound 3 (9.2 g) in methanol (115 mL) was carefully added to a solution of KOH in isopropanol (100 mL). A yellow potassium salt of the sulfonate was obtained in 85% yield after vacuum-drying for 12 hours. A portion of the 2,3,3-trimethylindolenium-5-sulfonate potassium salt (4 g) and 1,3-propanesultone (2.1 g) was heated in dichlorobenzene (40 mL) at 110 °C for 12 hours. The mixture was allowed to cool to room temperature and the resulting precipitate was filtered and washed with isopropanol. The resulting pink powder was dried under vacuum to give 97% of the desired compound.

Other compounds prepared by a similar method described above include polyhydroxyl indoles such as



## EXAMPLE 2

15

### Synthesis of indole disulfonate

(Figure 1, Compound 5,  $Y_7 = \text{SO}_3^-$ ;  $X_7 = \text{H}$ ;  $n = 2$ )

This compound was prepared by the same procedure described

in Example 1, except that 1,4-butanedisulfone was used in place of 1,3-propanedisulfone.

### EXAMPLE 3

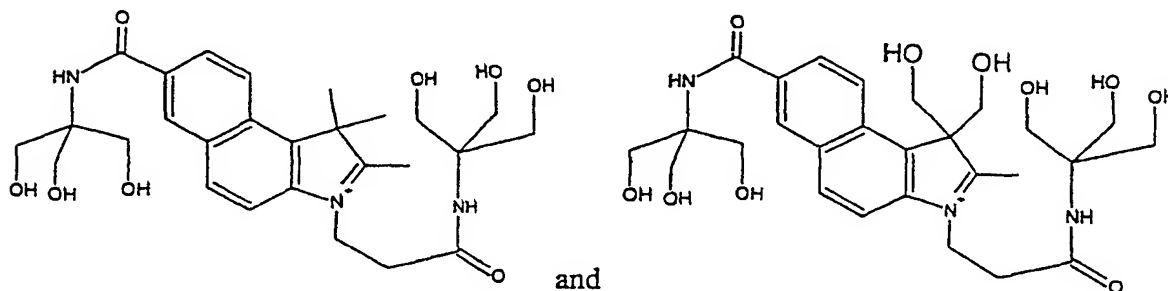
#### Synthesis of benzoindole disulfonate

5 (Figure 2, Compound 8,  $Y_7, Y_8 = SO_3^-$ ;  $X_7 = H$ ;  $n = 2$ )

This compound was prepared by the same procedure described in Example 1, except that hydrazinonaphthalenedisulfonic acid was used in place of hydrazinobenzenesulfonic acid.

Other compounds prepared by a similar method include

10 polyhydroxyindoles such as:



### EXAMPLE 4

#### Synthesis of benzoindole disulfonate

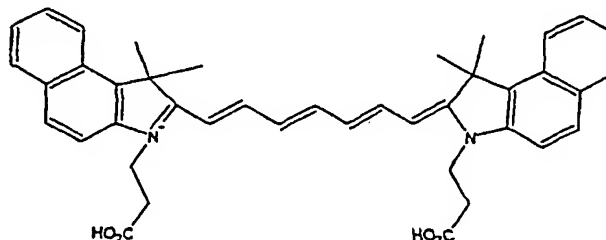
(Figure 2, Compound 8,  $Y_7, Y_8 = SO_3^-$ ;  $X_7 = OH$ ;  $n = 4$ )

This compound was prepared by the same procedure described  
15 in Example 1, except that 3-hydroxymethyl-4-hydroxyl-2-butanone was used

in place of 3-methyl-2-butanone.

### EXAMPLE 5

#### Synthesis of Bis(ethylcarboxymethyl)indocyanine Dye

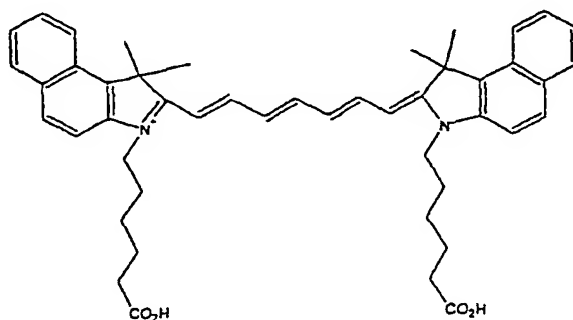


A mixture of 1,1,2-trimethyl-[1H]-benz[e]indole (9.1 g, 43.58  
5 mmoles) and 3-bromopropanoic acid (10.0 g, 65.37 mmoles) in 1,2-dichlorobenzene (40 mL) was heated at 110 °C for 12 hours. The solution was cooled to room temperature and the red residue obtained was filtered and washed with acetonitrile: diethyl ether (1:1) mixture. The solid obtained was dried under vacuum to give 10 g (64%) of light brown powder. A  
10 portion of this solid (6.0 g; 16.56 mmoles), glutaconaldehyde dianil monohydrochloride (2.36 g, 8.28 mmoles) and sodium acetate trihydrate (2.93 g, 21.53 mmoles) in ethanol (150 mL) were refluxed for 90 minutes. After evaporating the solvent, 40 mL of 2 N aqueous HCl was added to the residue and the mixture was centrifuged and the supernatant was decanted.  
15 This procedure was repeated until the supernatant became nearly colorless. About 5 mL of water:acetonitrile (3:2) mixture was added to the solid residue and lyophilized to obtain 2 g of dark green flakes. The purity of the

compound was established with  $^1\text{H}$ -NMR and liquid chromatography/mass spectrometry (LC/MS).

### EXAMPLE 6

### Synthesis of Bis(pentylcarboxymethyl)indocyanine Dye



5 A mixture of 2,2,3-trimethyl-[1H]-benz[e]indole (20 g, 95.6  
mmoles) and 6-bromohexanoic acid (28.1 g, 144.1 mmoles) in 1,2-  
dichlorobenzene (250 mL) was heated at 110 C for 12 hours. The green  
solution was cooled to room temperature and the brown solid precipitate  
formed was collected by filtration. After washing the solid with 1,2-  
10 dichlorobenzene and diethyl ether, the brown powder obtained (24 g, 64%)  
was dried under vacuum at room temperature. A portion of this solid (4.0 g;  
9.8 mmoles), glutaconaldehyde dianil monohydrochloride (1.4 g, 5 mmoles)  
and sodium acetate trihydrate (1.8 g, 12.9 mmoles) in ethanol (80 mL) were  
refluxed for 1 hour. After evaporating the solvent, 20 mL of a 2 N aqueous  
15 HCl was added to the residue and the mixture was centrifuged and the  
supernatant was decanted. This procedure was repeated until the

supernatant became nearly colorless. About 5 mL of water:acetonitrile (3:2) mixture was added to the solid residue and lyophilized to obtain about 2 g of dark green flakes. The purity of the compound was established with  $^1\text{H}$ -NMR, HPLC, and LC-MS.

5

### EXAMPLE 7

#### Synthesis of polyhydroxyindole sulfonate (Figure 3, Compound 13, $\text{Y}_7, \text{Y}_8 = \text{O}_3^-$ ; $\text{X}_7 = \text{OH}$ ; $n = 2$ )

Phosphorus oxychloride (37 ml, 0.4 mole) was added dropwise with stirring to a cooled ( $-2^\circ\text{C}$ ) mixture of dimethylformamide (DMF, 0.5 mole, 40 mL) and dichloromethane (DCM, 40 mL), followed by the addition of acetone (5.8 g, 0.1 mole). The ice bath was removed and the solution refluxed for 3 hours. After cooling to room temperature, the product was either partitioned in water/DCM, separated and dried, or was purified by fractional distillation. Nuclear magnetic resonance and mass spectral analyses showed that the desired intermediate, **10**, was obtained. Reaction of the intermediate with 2 equivalents of 2,2,3-trimethyl-[H]-benz[e]indolesulfonate-N-propanoic acid and 2 equivalents of sodium acetate trihydrate in ethanol gave a blue-green solution after 1.5 hours at reflux. Further functionalization of the dye with bis(isopropylidene)acetal protected monosaccharide is effected by the method described in the literature (J. H. Flanagan, C. V. Owens, S. E. Romero, et al., Near infrared heavy-atom-modified fluorescent dyes for base-calling in DNA-sequencing application using temporal discrimination. *Anal. Chem.*, 1998, 70(13), 2676-2684).

### EXAMPLE 8

Synthesis of polyhydroxyindole sulfonate  
(Figure 4, Compound 16,  $Y_7, Y_8 = SO_3^-$ ;  $X_7 = H$ ;  $n = 1$ )

Preparation of this compound was readily accomplished by the same procedure described in Example 6 using p-hydroxybenzenesulfonic acid in the place of the monosaccharide, and benzoindole instead of indole derivatives.

### EXAMPLE 9

Synthesis of polyhydroxyindole sulfonate  
(Figure 5, Compound 20,  $Y_7, Y_8 = H$ ;  $X_7 = OH$ ;  $n = 1$ )

The hydroxyindole compound was readily prepared by a literature method (P.L. Southwick, J.G. Cairns, L.A. Ernst, and A.S. Waggoner, One pot Fischer synthesis of (2,3,3-trimethyl-3-H-indol-5-yl)-acetic acid derivatives as intermediates for fluorescent biolabels. *Org. Prep. Proced. Int. Briefs*, 1988, 20(3), 279-284). Reaction of p-carboxymethylphenylhydrazine hydrochloride (30 mmol, 1 equiv.) and 1,1-bis(hydroxymethyl)propanone (45 mmol, 1.5 equiv.) in acetic acid (50 mL) at room temperature for 30 minutes and at reflux for 1 gave (3,3-dihydroxymethyl-2-methyl-3-H-indol-5-yl)-acetic acid as a solid residue.

The intermediate 2-chloro-1-formyl-3-hydroxymethylenecyclohexane was prepared as described in the literature (G. A. Reynolds and K. H. Drexhage, Stable heptamethine pyrylium dyes that absorb in the infrared. *J.*



Org. Chem., 1977, 42(5), 885-888). Equal volumes (40 mL each) of dimethylformamide (DMF) and dichloromethane were mixed and the solution was cooled to -10°C in acetone-dry ice bath. Under argon atmosphere, phosphorus oxychloride (40 mL) in dichloromethane was added dropwise to the cool DMF solution, followed by the addition of 10 g of cyclohexanone. The resulting solution was allowed to warm up to room temperature and heated at reflux for 6 hours. After cooling to room temperature, the mixture was poured into ice-cold water and stored at 4°C for 12 hours. A yellow powder was obtained. Condensation of a portion of this cyclic dialdehyde (1 equivalent) with the indole intermediate (2 equivalents) was carried out as described in Example 5. Further, the functionalization of the dye with bis (isopropylidene)acetal protected monosaccharide was effected by the method described in the literature (J. H. Flanagan, C. V. Owens, S. E. Romero, et al., Near infrared heavy-atom-modified fluorescent dyes for base-calling in DNA-sequencing application using temporal discrimination. Anal. Chem., 1998, 70(13), 2676-2684).

#### EXAMPLE 10

Synthesis of polyhydroxybenzoindole sulfonate  
(Figure 6, Compound 22, Y<sub>7</sub>, Y<sub>8</sub>=H; X<sub>7</sub>=OH; n=1)

A similar method described in Example 8 was used to prepare this compound by replacing the indole with benzoindole derivatives.

EXAMPLE 11Synthesis of rigid heteroatomic indole sulfonate  
(Figure 7, Compound 27, Y<sub>7</sub>, Y<sub>8</sub>, X<sub>7</sub> = H; n = 1)

Starting with 3-oxo-4-cyclohexenone, this heteroatomic  
5 hydrophilic dye was readily prepared as described in Example 8.

EXAMPLE 12Minimally invasive monitoring of the blood clearance profile of the dyes

A laser of appropriate wavelength for excitation of the dye  
chromophore was directed into one end of a fiber optic bundle and the other  
10 end was positioned a few millimeters from the ear of a rat. A second fiber  
optic bundle was also positioned near the same ear to detect the emitted  
fluorescent light, and the other end was directed into the optics and  
electronics for data collection. An interference filter (IF) in the collection  
optics train was used to select emitted fluorescent light of the appropriate  
15 wavelength for the dye chromophore.

Sprague-Dawley or Fischer 344 rats were anesthetized with  
urethane administered via intraperitoneal injection at a dose of 1.35 g/kg  
body weight. After the animals had achieved the desired plane of  
anesthesia, a 21 gauge butterfly with 12" tubing was placed in the lateral tail  
20 vein of each animal and flushed with heparinized saline. The animals were  
placed onto a heating pad and kept warm throughout the entire study. The  
lobe of the left ear was affixed to a glass microscope slide to reduce  
movement and vibration.

Incident laser light delivered from the fiber optic was centered on the affixed ear. Data acquisition was then initiated, and a background reading of fluorescence was obtained prior to administration of the test agent.

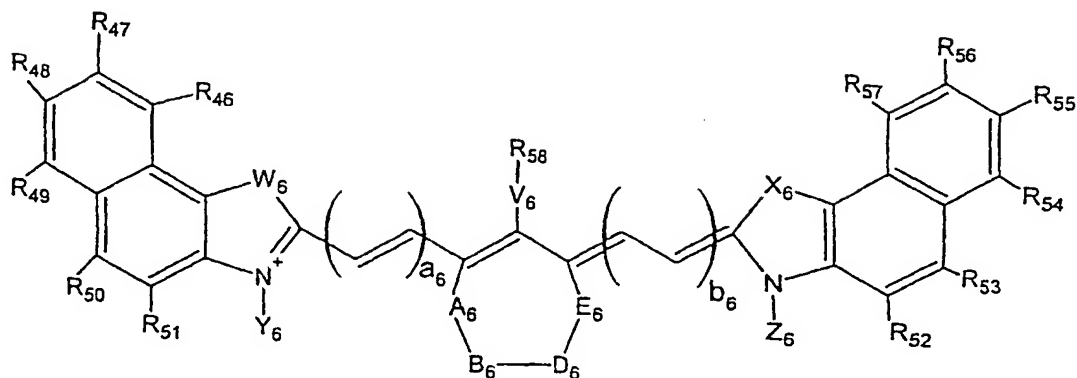
5                   The compound was administered to the animal through a bolus injection in the lateral tail vein. The dose was typically 0.05 to 20  $\mu$ mole/kg of body weight. The fluorescence signal rapidly increased to a peak value, then decayed as a function of time as the conjugate cleared from the bloodstream.

10                   This procedure was repeated with several dye-peptide conjugates in normal and tumored rats. Representative profiles are shown in Figures 6-10.

                  While the invention has been disclosed by reference to the details of preferred embodiments of the invention, it is to be understood that  
15                   the disclosure is intended in an illustrative rather than in a limiting sense, as it is contemplated that modifications will readily occur to those skilled in the art, within the spirit of the invention and the scope of the appended claims.

What is claimed is:

1. A composition comprising benzoindeole of formula



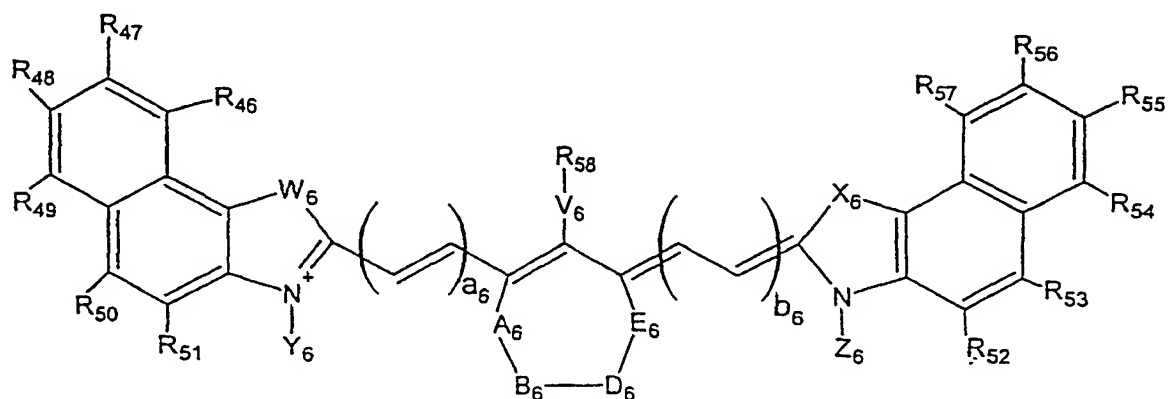
- wherein  $R_{46}$ ,  $R_{47}$ ,  $R_{48}$ ,  $R_{49}$ ,  $R_{50}$ ,  $R_{51}$ ,  $R_{52}$ ,  $R_{53}$ ,  $R_{54}$ ,  $R_{55}$ ,  $R_{56}$ ,  $R_{57}$  and  $R_{58}$ ,  $Y_6$ , and  $Z_6$  are independently selected from the group consisting of -H, C1-C10 alkoxy, C1-  
 5 C10 polyalkoxyalkyl, C1-C20 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, saccharides, amino, C1-C10 aminoalkyl, cyano, nitro, halogen, hydrophilic peptides, arylpolysulfonates, C1-C10 alkyl, C5-C20 aryl,  $-SO_3T$ ,  $-CO_2T$ ,  $-OH$ ,  $-(CH_2)_aSO_3T$ ,  $-(CH_2)_aOSO_3T$ ,  $-(CH_2)_aNHCO_2(CH_2)_bSO_3T$ , -  
 $(CH_2)_aOCO(CH_2)_bSO_3T$ ,  $-(CH_2)_aCONH(CH_2)_bSO_3T$ ,  $-(CH_2)_aNHCO(CH_2)_bSO_3T$ , -  
 10  $(CH_2)_aNHCONH(CH_2)_bSO_3T$ ,  $-(CH_2)_aNHCSNH(CH_2)_bSO_3T$ , -  
 $(CH_2)_aOCONH(CH_2)_bSO_3T$ ,  $-(CH_2)_aPO_3HT$ ,  $-(CH_2)_aPO_3T_2$ ,  $-(CH_2)_aOPO_3HT$ , -  
 $(CH_2)_aOPO_3T_2$ ,  $-(CH_2)_aNHPO_3HT$ ,  $-(CH_2)_aNHPO_3T_2$ ,  $-(CH_2)_aCO_2(CH_2)_bPO_3HT$ , -

- $(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 $(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{CONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 15  $(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCO}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 $(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 $(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ ,  $-(\text{CH}_2)_a\text{NHCSNH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ , -  
 $(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{PO}_3\text{HT}$ , and  $-(\text{CH}_2)_a\text{OCONH}(\text{CH}_2)_b\text{PO}_3\text{T}_2$ ,  $-\text{CH}_2(\text{CH}_2\text{-O-}$   
 $\text{CH}_2)_c\text{-CH}_2\text{-OH}$ ,  $-(\text{CH}_2)_d\text{-CO}_2\text{T}$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_e\text{-CH}_2\text{-CO}_2\text{T}$ ,  $-(\text{CH}_2)_f\text{-NH}_2$ ,  $-\text{CH}_2\text{-}$   
 20  $(\text{CH}_2\text{-O-CH}_2)_g\text{-CH}_2\text{-NH}_2$ ,  $-(\text{CH}_2)_h\text{-N(R}_a\text{)-(CH}_2)_i\text{-CO}_2\text{T}$ , and  $-(\text{CH}_2)_j\text{-N(R}_b\text{)-CH}_2\text{-(CH}_2\text{-}$   
 $\text{O-CH}_2)_k\text{-CH}_2\text{-CO}_2\text{T}$ ;  $W_6$  and  $X_6$  are selected from the group consisting of  
 $-\text{CR}_c\text{R}_d$ ,  $-\text{O-}$ ,  $-\text{NR}_c$ ,  $-\text{S-}$ , and  $-\text{Se-}$ ;  $V_6$  is a single bond or is selected from the  
group consisting of  $-\text{O-}$ ,  $-\text{S-}$ ,  $-\text{Se-}$ , and  $-\text{NR}_a$ ;  $D_6$  is a single or a double bond;  
 $A_6$ ,  $B_6$  and  $E_6$  may be the same or different and are selected from the group  
 25 consisting of  $-\text{O-}$ ,  $-\text{S-}$ ,  $-\text{Se-}$ ,  $-\text{P-}$ ,  $-\text{NR}_a$ ,  $-\text{CR}_c\text{R}_d$ ,  $\text{CR}_c$ , alkyl, and  $-\text{C=O}$ ;  $A_6$ ,  $B_6$ ,  
 $D_6$ , and  $E_6$  may together form a 6 or 7 membered carbocyclic ring or a 6 or 7  
membered heterocyclic ring optionally containing one or more oxygen,  
nitrogen, or sulfur atoms;  $a$ ,  $b$ ,  $d$ ,  $f$ ,  $h$ ,  $i$ , and  $j$  independently vary from 1-10;  
 $c$ ,  $e$ ,  $g$ , and  $k$  independently vary from 1-100;  $a_6$  and  $b_6$  vary from 0 to 5;  $R_a$ ,  
 30  $R_b$ ,  $R_c$ , and  $R_d$  are defined in the same manner as  $Y_6$ ;  $T$  is either H or a  
negative charge.

2. The composition of claim 1 wherein  $R_{46}$ ,  $R_{47}$ ,  $R_{48}$ ,  $R_{49}$ ,  $R_{50}$ ,  $R_{51}$ ,  $R_{52}$ ,  $R_{53}$ ,  $R_{54}$ ,  $R_{55}$ ,  $R_{56}$ ,  $R_{57}$ , and  $R_{58}$ ,  $Y_6$ , and  $Z_6$  are independently selected from the group consisting of -H, C1-C5 alkoxyl, C1-C5 polyalkoxyalkyl, C1-C10 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, mono- and disaccharides, nitro, hydrophilic peptides, arylpolysulfonates, C1-C5 alkyl, C5-C20 aryl,  $-\text{SO}_3\text{T}$ ,  $-\text{CO}_2\text{T}$ ,  $-\text{OH}$ ,  $-(\text{CH}_2)_a\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-\text{CH}_2(\text{CH}_2\text{-O-CH}_2)_c\text{-CH}_2\text{-OH}$ ,  $-(\text{CH}_2)_d\text{-CO}_2\text{T}$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_e\text{-CH}_2\text{-CO}_2\text{T}$ ,  $-(\text{CH}_2)_f\text{-NH}_2$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_g\text{-CH}_2\text{-NH}_2$ ,  $-(\text{CH}_2)_h\text{-N(R}_a\text{)-(CH}_2)_i\text{-CO}_2\text{T}$ , and  $-(\text{CH}_2)_j\text{-N(R}_b\text{)-CH}_2\text{-(CH}_2\text{-O-CH}_2)_k\text{-CH}_2\text{-CO}_2\text{T}$ ;  $W_6$  and  $X_6$  are selected from the group consisting of  $-\text{CR}_c\text{R}_d$ ,  $-\text{O-}$ ,  $-\text{NR}_c$ ,  $-\text{S-}$ , and  $-\text{Se-}$ ;  $V_6$  is a single bond or is selected from the group consisting of  $-\text{O-}$ ,  $-\text{S-}$ ,  $-\text{Se-}$ , and  $-\text{NR}_a$ ;  $D_6$  is a single or a double bond;  $A_6$ ,  $B_6$  and  $E_6$  are independently selected from the group consisting of  $-\text{O-}$ ,  $-\text{S-}$ ,  $-\text{NR}_a$ ,  $-\text{CR}_c\text{R}_d$ ,  $\text{CR}_c$ , and alkyl;  $A_6$ ,  $B_6$ ,  $D_6$ , and  $E_6$  may together form a 6 or 7 membered carbocyclic ring or a 6 or 7 membered heterocyclic ring optionally containing one or two oxygen, nitrogen, or sulfur atom;  $a$ ,  $b$ ,  $d$ ,  $f$ ,  $h$ ,  $i$ , and  $j$  independently vary from 1-5;  $c$ ,  $e$ ,  $g$ , and  $k$  independently vary from 1-50;  $a_5$  and  $b_5$  independently vary from 0 to 5;  $R_a$ ,  $R_b$ ,  $R_c$ , and  $R_d$  are defined in the same manner as  $Y_6$ ; and  $T$  is a negative charge.

3. The composition of claim 1 wherein each of  $R_{46}$ ,  $R_{48}$ ,  $R_{50}$ ,  $R_{51}$ ,  $R_{52}$  and  $R_{53}$ ,  $R_{55}$ , and  $R_{57}$  is H; each of  $R_{47}$ ,  $R_{49}$ ,  $R_{54}$ , and  $R_{56}$  is  $\text{SO}_3\text{T}$ ;  $R_{58}$  is glucose; each of  $Y_5$  and  $Z_6$  is  $-(\text{CH}_2)_3\text{SO}_3\text{T}$ ; each of  $W_6$  and  $X_6$  is  $-\text{C}(\text{CH}_2\text{OH})_2$ ;  $D_6$  is a double bond; each of  $B_6$  and  $E_6$  is  $-\text{CH}-$ ;  $A_6$  is  $-\text{CH}_2-$ ; each of  $a_6$  and  $b_6$  is 1;
- 5  $V_5$  is a single bond; and T is a negative charge.

4. A method for performing a diagnostic or therapeutic procedure comprising administering to a mammal an effective amount of the benzoindole of formula



- 5 wherein  $R_{46}$ ,  $R_{47}$ ,  $R_{48}$ ,  $R_{49}$ ,  $R_{50}$ ,  $R_{51}$ ,  $R_{52}$ ,  $R_{53}$ ,  $R_{54}$ ,  $R_{55}$ ,  $R_{56}$ ,  $R_{57}$ , and  $R_{58}$ ,  $Y_6$ , and  $Z_6$  are independently selected from the group consisting of -H, C1-C10 alkoxy, C1-C10 polyalkoxyalkyl, C1-C20 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, saccharides, amino, C1-C10 aminoalkyl, cyano, nitro, halogen, hydrophilic peptides, arylpolysulfonates, C1-C10 alkyl, C5-C20 aryl,  $-SO_3T$ ,  $-CO_2T$ ,  $-OH$ ,  $-(CH_2)_aSO_3T$ ,  $-(CH_2)_aOSO_3T$ ,  $-(CH_2)_aNHSO_3T$ ,  $-(CH_2)_aCO_2(CH_2)_bSO_3T$ ,  $-(CH_2)_aOCO(CH_2)_bSO_3T$ ,  $-(CH_2)_aCONH(CH_2)_bSO_3T$ ,  $-(CH_2)_aNHCO(CH_2)_bSO_3T$ ,  $-(CH_2)_aNHCONH(CH_2)_bSO_3T$ ,  $-(CH_2)_aNHCSNH(CH_2)_bSO_3T$ , -
- 10



- $(CH_2)_aOCONH(CH_2)_bSO_3T$ ,  $-(CH_2)_aPO_3HT$ ,  $-(CH_2)_aPO_3T_2$ ,  $-(CH_2)_aOPO_3HT$ , -  
 $(CH_2)_aOPO_3T_2$ ,  $-(CH_2)_aNHPO_3HT$ ,  $-(CH_2)_aNHPO_3T_2$ ,  $-(CH_2)_aCO_2(CH_2)_bPO_3HT$ , -  
 15  $(CH_2)_aCO_2(CH_2)_bPO_3T_2$ ,  $-(CH_2)_aOCO(CH_2)_bPO_3HT$ ,  $-(CH_2)_aOCO(CH_2)_bPO_3T_2$ , -  
 $(CH_2)_aCONH(CH_2)_bPO_3HT$ ,  $-(CH_2)_aCONH(CH_2)_bPO_3T_2$ , -  
 $(CH_2)_aNHCO(CH_2)_bPO_3HT$ ,  $-(CH_2)_aNHCO(CH_2)_bPO_3T_2$ , -  
 $(CH_2)_aNHCONH(CH_2)_bPO_3HT$ ,  $-(CH_2)_aNHCONH(CH_2)_bPO_3T_2$ , -  
 $(CH_2)_aNHCSNH(CH_2)_bPO_3HT$ ,  $-(CH_2)_aNHCSNH(CH_2)_bPO_3T_2$ , -  
 20  $(CH_2)_aOCONH(CH_2)_bPO_3HT$ , and  $-(CH_2)_aOCONH(CH_2)_bPO_3T_2$ ,  $-CH_2(CH_2-O-$   
 $CH_2)_c-CH_2-OH$ ,  $-(CH_2)_d-CO_2T$ ,  $-CH_2-(CH_2-O-CH_2)_e-CH_2-CO_2T$ ,  $-(CH_2)_f-NH_2$ ,  $-CH_2-$   
 $(CH_2-O-CH_2)_g-CH_2-NH_2$ ,  $-(CH_2)_h-N(R_a)-(CH_2)_i-CO_2T$ , and  $-(CH_2)_j-N(R_b)-CH_2-(CH_2-$   
 $O-CH_2)_k-CH_2-CO_2T$ ;  $W_6$  and  $X_6$  are selected from the group consisting of  
 $-CR_cR_d$ ,  $-O-$ ,  $-NR_c$ ,  $-S-$ , and  $-Se$ ;  $V_6$  is a single bond or is selected from the  
 25 group consisting of  $-O-$ ,  $-S-$ ,  $-Se-$ , and  $-NR_a$ ;  $D_6$  is a single or a double bond;  
 $A_6$ ,  $B_6$  and  $E_6$  may be the same or different and are selected from the group  
 consisting of  $-O-$ ,  $-S-$ ,  $-Se-$ ,  $-P-$ ,  $-NR_a$ ,  $-CR_cR_d$ ,  $CR_c$ , alkyl, and  $-C=O$ ;  $A_6$ ,  $B_6$ ,  
 $D_6$ , and  $E_6$  may together form a 6 or 7 membered carbocyclic ring or a 6 or 7  
 membered heterocyclic ring optionally containing one or more oxygen,  
 30 nitrogen, or sulfur atom;  $a$ ,  $b$ ,  $d$ ,  $f$ ,  $h$ ,  $i$ , and  $j$  independently vary from 1-10;  
 $c$ ,  $e$ ,  $g$ , and  $k$  independently vary from 1-100;  $a_6$  and  $b_6$  vary from 0 to 5;  $R_a$ ,  
 $R_b$ ,  $R_c$ , and  $R_d$  are defined in the same manner as  $Y_6$ ; and  $T$  is either H or a  
 negative charge.

5. The method of claim 4 comprising administering an effective amount of the composition of indoles wherein  $R_{46}$ ,  $R_{47}$ ,  $R_{48}$ ,  $R_{49}$ ,  $R_{50}$ ,  $R_{51}$ ,  $R_{52}$ ,  $R_{53}$ ,  $R_{54}$ ,  $R_{55}$ ,  $R_{56}$ ,  $R_{57}$ , and  $R_{58}$ ,  $Y_6$ , and  $Z_6$  are independently selected from the group consisting of -H, C1-C5 alkoxy, C1-C5 polyalkoxyalkyl, C1-C10 polyhydroxyalkyl, C5-C20 polyhydroxyaryl, mono- and disaccharides, nitro, hydrophilic peptides, arylpolysulfonates, C1-C5 alkyl, C5-C20 aryl,  $-\text{SO}_3\text{T}$ ,  $-\text{CO}_2\text{T}$ ,  $-\text{OH}$ ,  $-(\text{CH}_2)_a\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{NHSO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{CO}_2(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-(\text{CH}_2)_a\text{OCO}(\text{CH}_2)_b\text{SO}_3\text{T}$ ,  $-\text{CH}_2(\text{CH}_2\text{-O-CH}_2)_c\text{-CH}_2\text{-OH}$ ,  $-(\text{CH}_2)_d\text{-CO}_2\text{T}$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_e\text{-CH}_2\text{-CO}_2\text{T}$ ,  $-(\text{CH}_2)_f\text{-NH}_2$ ,  $-\text{CH}_2\text{-(CH}_2\text{-O-CH}_2)_g\text{-CH}_2\text{-NH}_2$ ,  $-(\text{CH}_2)_h\text{-N(R}_a\text{)-(CH}_2)_i\text{-CO}_2\text{T}$ , and  $-(\text{CH}_2)_j\text{-N(R}_b\text{)-CH}_2\text{-(CH}_2\text{-O-CH}_2)_k\text{-CH}_2\text{-CO}_2\text{T}$ ;  $W_6$  and  $X_6$  are selected from the group consisting of  $-\text{CR}_c\text{R}_d$ ,  $-\text{O-}$ ,  $-\text{NR}_c$ ,  $-\text{S-}$ , and  $-\text{Se-}$ ;  $V_6$  is a single bond or is selected from the group consisting of  $-\text{O-}$ ,  $-\text{S-}$ ,  $-\text{Se-}$ , and  $-\text{NR}_a$ ;  $D_6$  is a single or a double bond;  $A_6$ ,  $B_6$  and  $E_6$  are independently selected from the group consisting of  $-\text{O-}$ ,  $-\text{S-}$ ,  $-\text{NR}_a$ ,  $-\text{CR}_c\text{R}_d$ ,  $\text{CR}_c$ , and alkyl;  $A_6$ ,  $B_6$ ,  $D_6$ , and  $E_6$  may together form a 6 or 7 membered carbocyclic ring or a 6 or 7 membered heterocyclic ring optionally containing one or two oxygen, nitrogen, or sulfur atom;  $a$ ,  $b$ ,  $d$ ,  $f$ ,  $h$ ,  $i$ , and  $j$  independently vary from 1-5;  $c$ ,  $e$ ,  $g$ , and  $k$  independently vary from 1-50;  $a_5$  and  $b_5$  independently vary from 0 to 5;  $R_a$ ,  $R_b$ ,  $R_c$ , and  $R_d$  are defined in the same manner as  $Y_6$ ; and  $T$  is a negative charge.

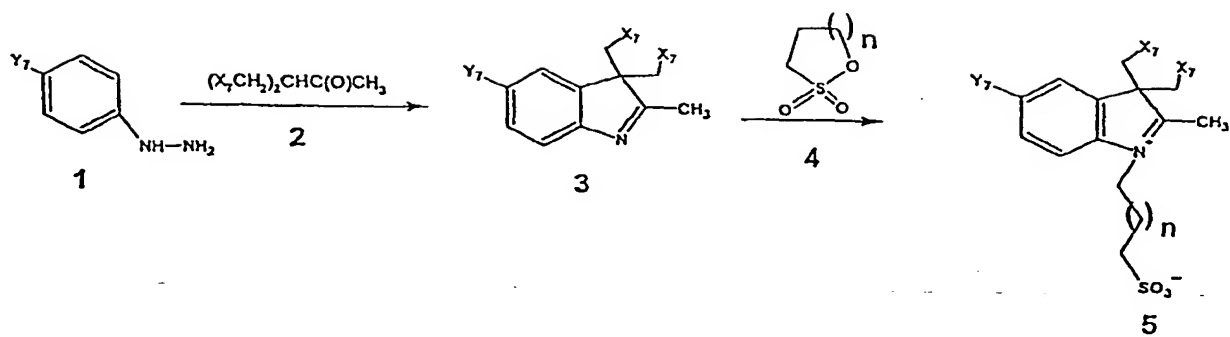
6. The method of claim 5 comprising administering an effective amount of the composition of indoles wherein each of  $R_{46}$ ,  $R_{48}$ ,  $R_{50}$  to  $R_{53}$ ,  $R_{55}$ , and  $R_{57}$  is H; each of  $R_{47}$ ,  $R_{49}$ ,  $R_{54}$ , and  $R_{56}$  is  $\text{SO}_3\text{T}$ ;  $R_{58}$  is glucose; each of  $Y_5$  and  $Z_6$  is  $-(\text{CH}_2)_3\text{SO}_3\text{T}$ ; each of  $W_6$  and  $X_6$  is  $-\text{C}(\text{CH}_2\text{OH})_2$ ;  $D_6$  is a double bond;
- 5 each of  $B_6$  and  $E_6$  is  $-\text{CH}-$ ;  $A_6$  is  $-\text{CH}_2-$ ; each of  $a_6$  and  $b_6$  is 1;  $V_5$  is a single bond; and T is a negative charge.
7. The method of claim 4 wherein said procedure utilizes light of wavelength in the region of 350-1300 nm.
8. The method of claim 4 wherein said procedure comprises monitoring a blood clearance profile by fluorescence using light of wavelength in the region of 350 to 1300 nm.
9. The method of claim 4 wherein said procedure comprises monitoring a blood clearance profile by absorption using light of wavelength in the region of 350 to 1300 nm.
10. The method of claim 4 wherein said procedure is for physiological function monitoring.
11. The method of claim 10 wherein said procedure is for renal function monitoring.

12. The method of claim 10 wherein said procedure is for cardiac function monitoring.

13. The method of claim 10 wherein said procedure is for kidney function monitoring.

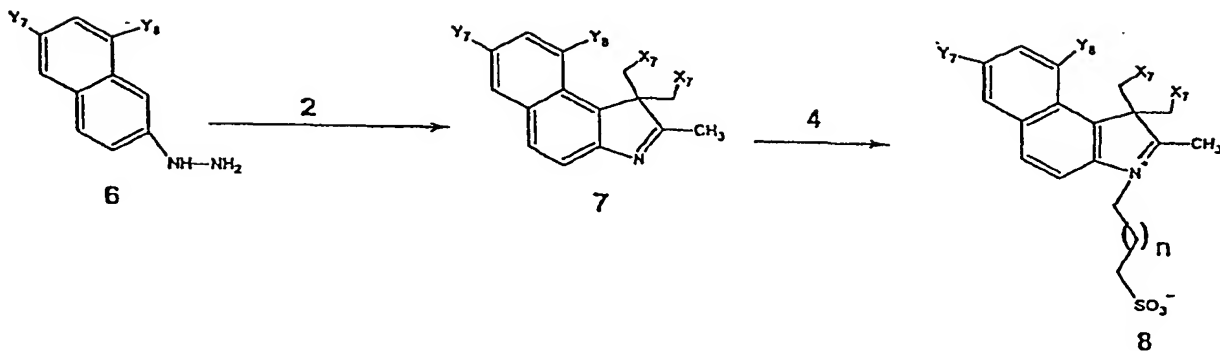
14. The method of claim 10 wherein said procedure is for determining organ perfusion *in vivo*.

FIGURE 1



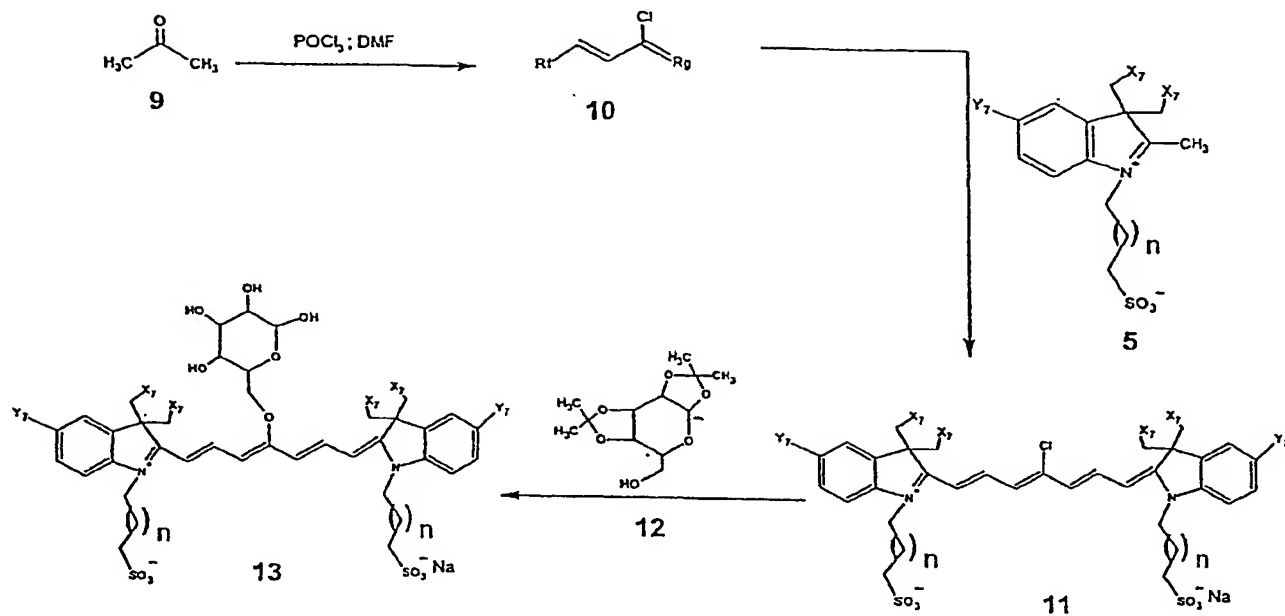
$n = 1-3$ ;  $X_7 = H, OH$ ;  $Y_7 = H, SO_3^-, CO_2H, CH_2CO_2H, CH_2OH$

FIGURE 2



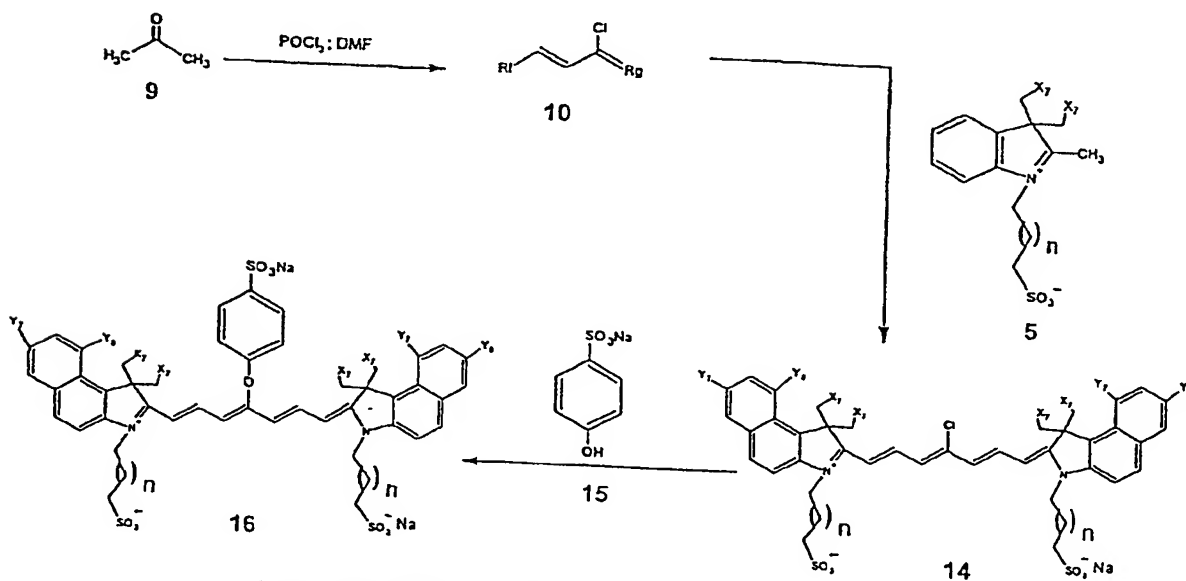
$n = 1-3$ ;  $X_7 = H, OH$ ;  $Y_7, Y_8 = H, SO_3^-, CO_2H, CH_2CO_2H, CH_2OH$

FIGURE 3



$n = 1-3$ ;  $\text{X}_7 = \text{H}, \text{OH}$ ;  $\text{Y}_7 = \text{H}, \text{SO}_3^-, \text{CO}_2\text{H}, \text{CH}_2\text{CO}_2\text{H}, \text{CH}_2\text{OH}$ ;  $\text{R}_1 = (\text{CH}_3)_2\text{N}$  or  $\text{OH}$ ;  $\text{R}_9 = (\text{CH}_3)_2\text{N}^+$  or  $\text{CHO}$

FIGURE 4



$n = 1-3$ ;  $\text{X}_7 = \text{H}, \text{OH}$ ;  $\text{Y}_7 = \text{H}, \text{SO}_3^-, \text{CO}_2\text{H}, \text{CH}_2\text{CO}_2\text{H}, \text{CH}_2\text{OH}$ ;  $\text{R}_1 = (\text{CH}_3)_2\text{N}$  or  $\text{OH}$ ;  $\text{R}_9 = (\text{CH}_3)_2\text{N}^+$  or  $\text{CHO}$

FIGURE 5

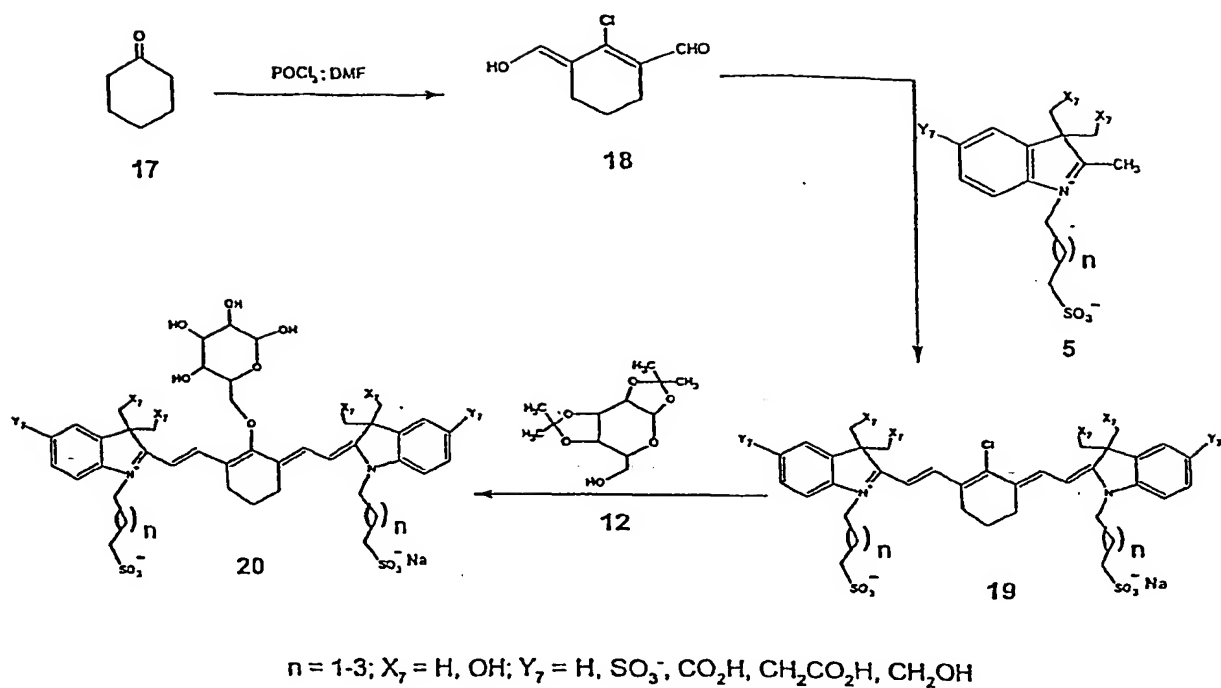


FIGURE 6

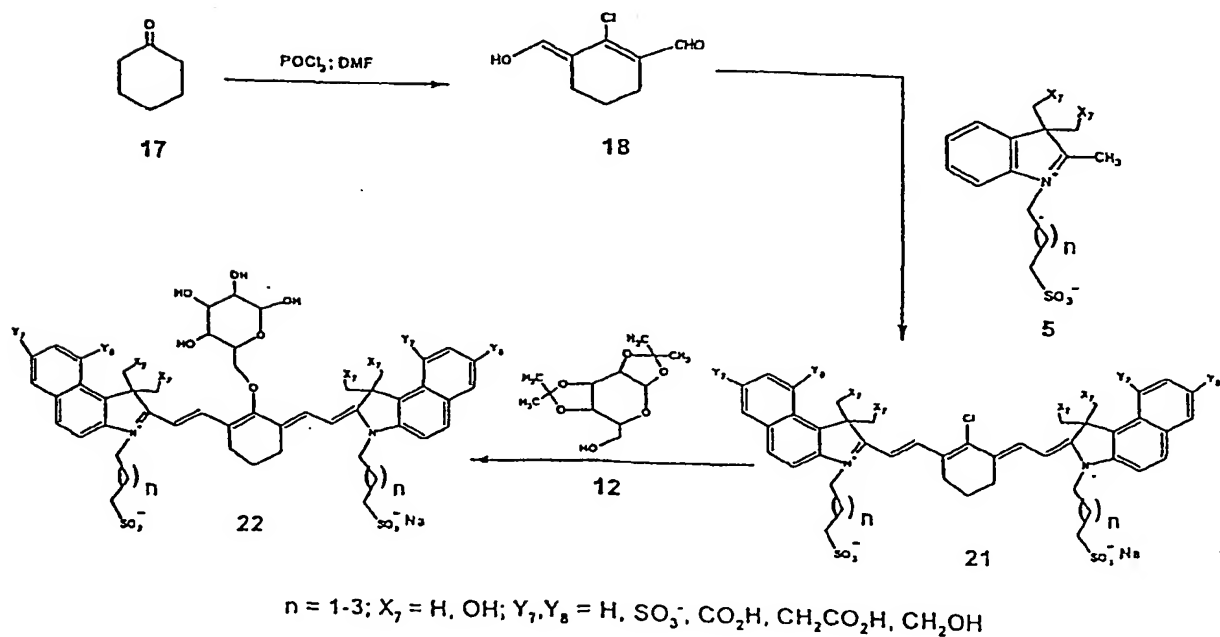
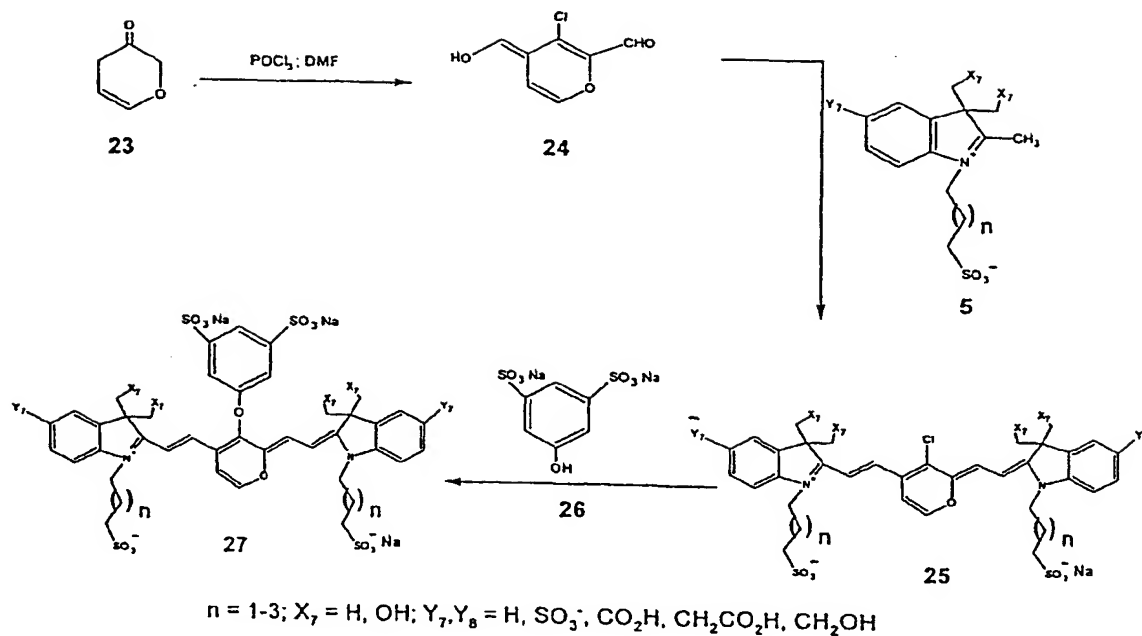


FIGURE 7





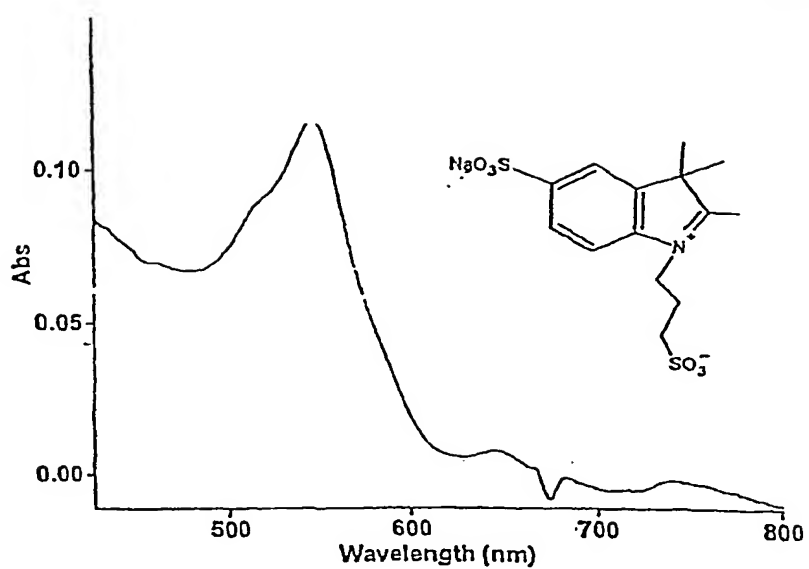


Figure 8a

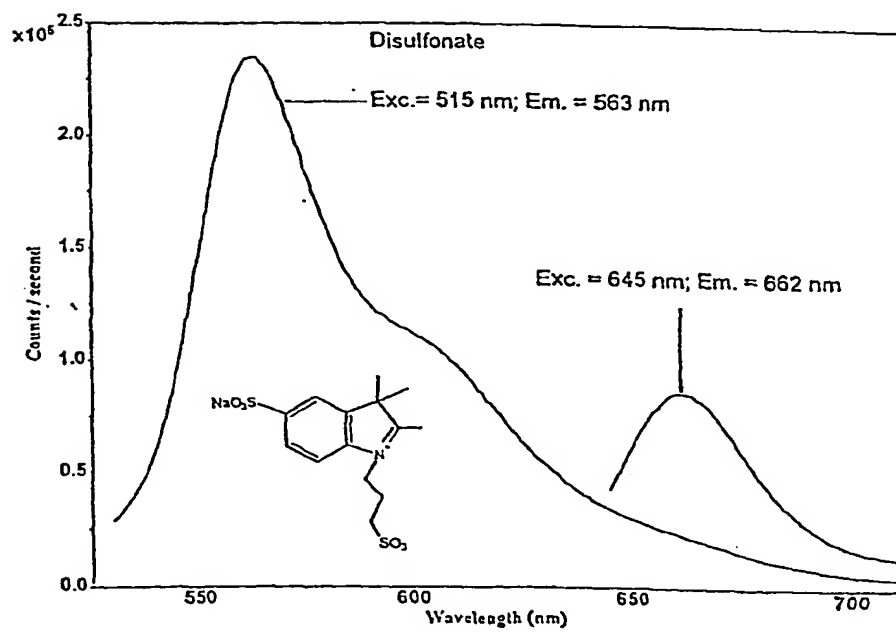


Figure 8b

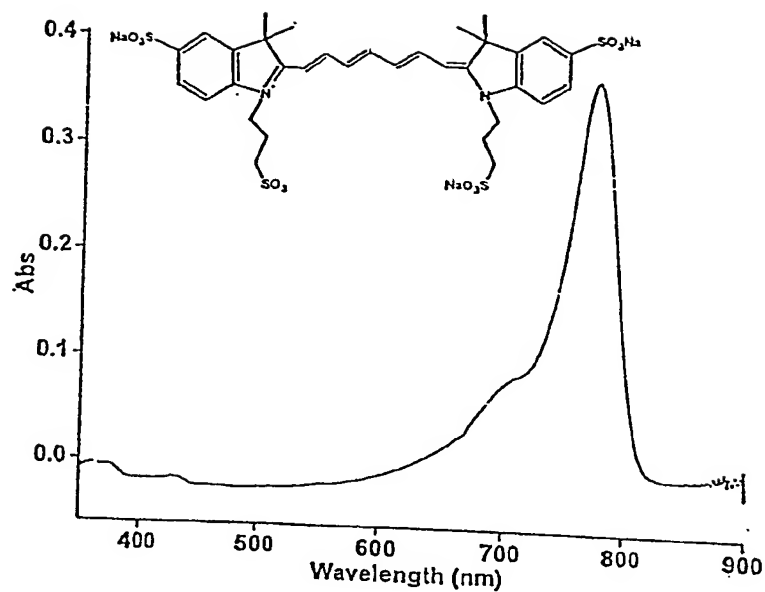


Figure 9a

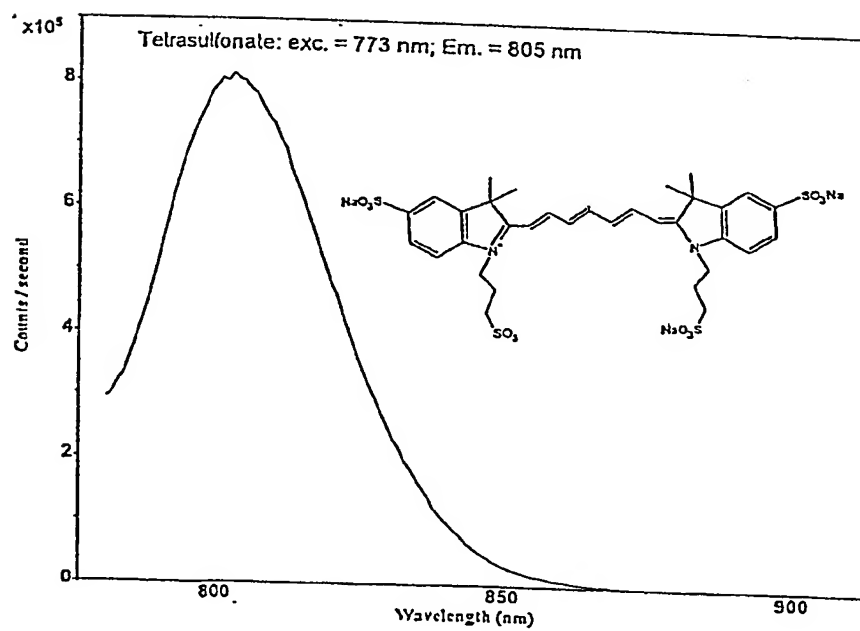


Figure 9b

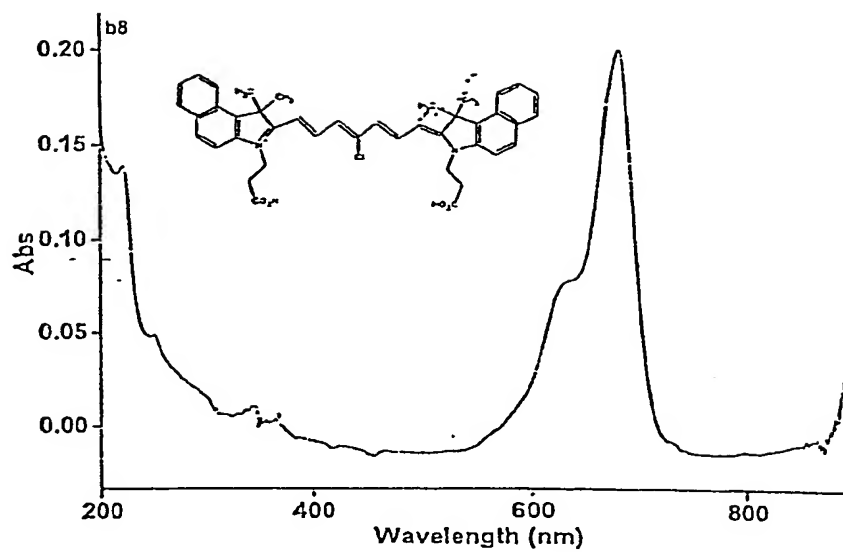


Figure 10a

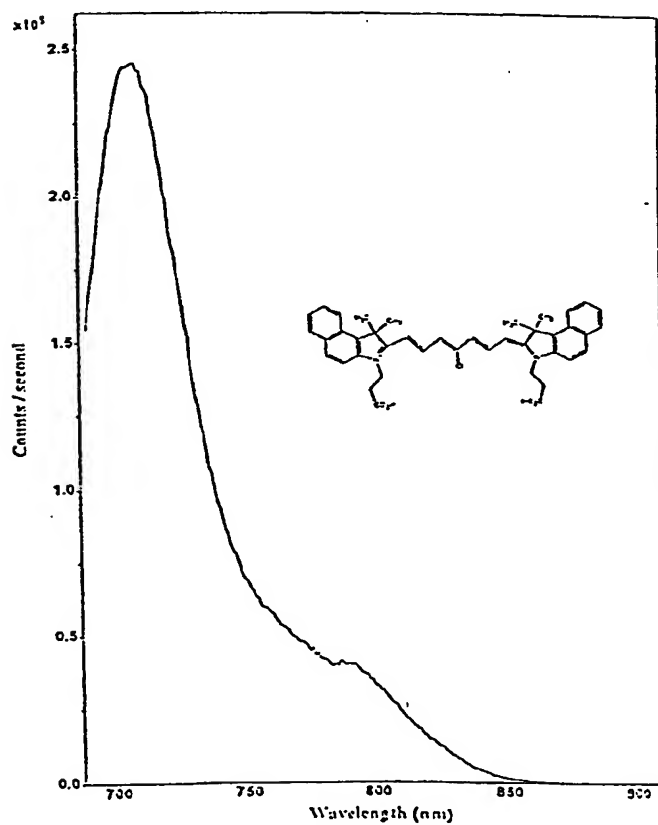


Figure 10b

# Blood clearance of hydrophilic polyaspartic acid-cyanine dye

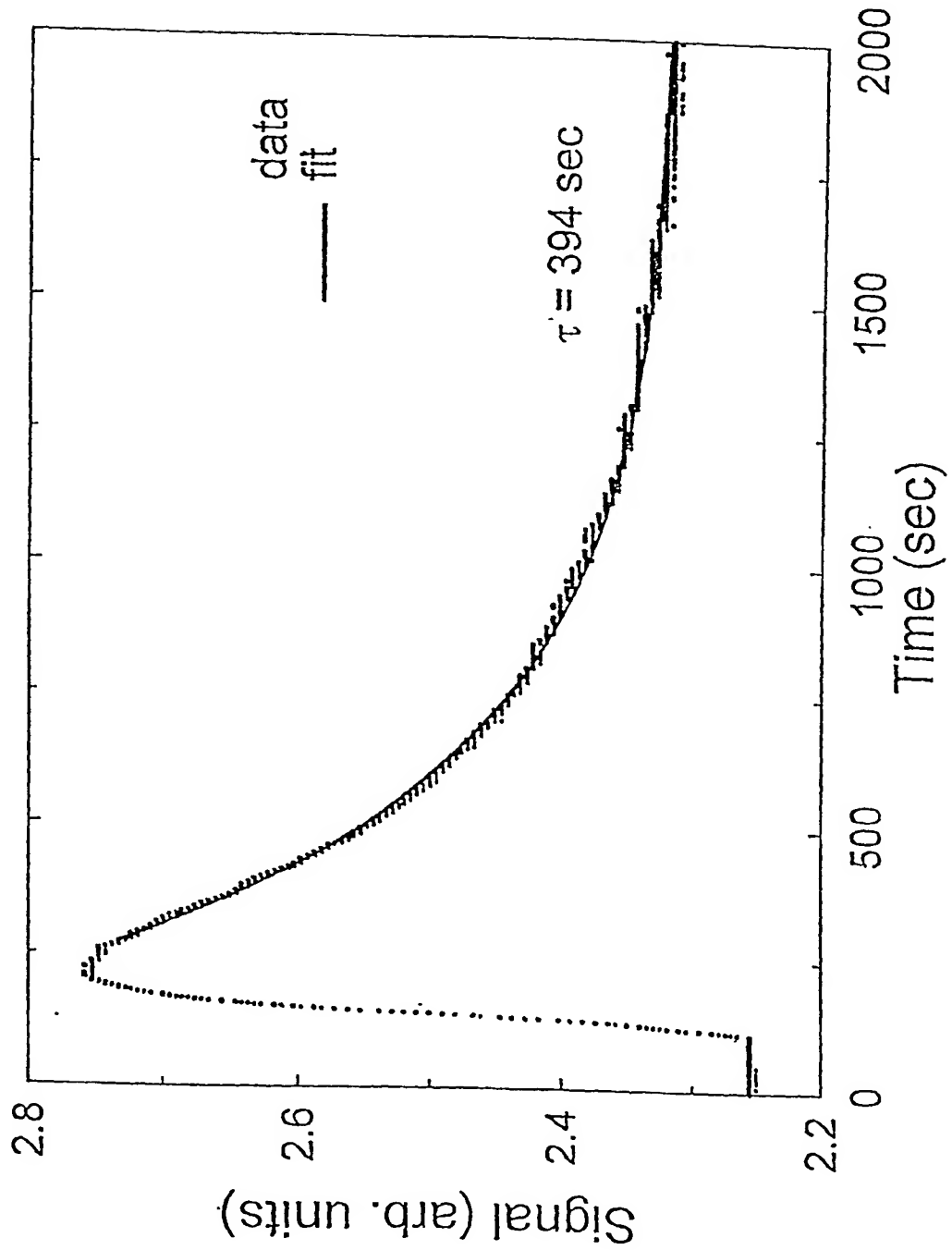


Figure 11

## Blood clearance profile of cyanine dye-polyaspartic acid (30 kDa)

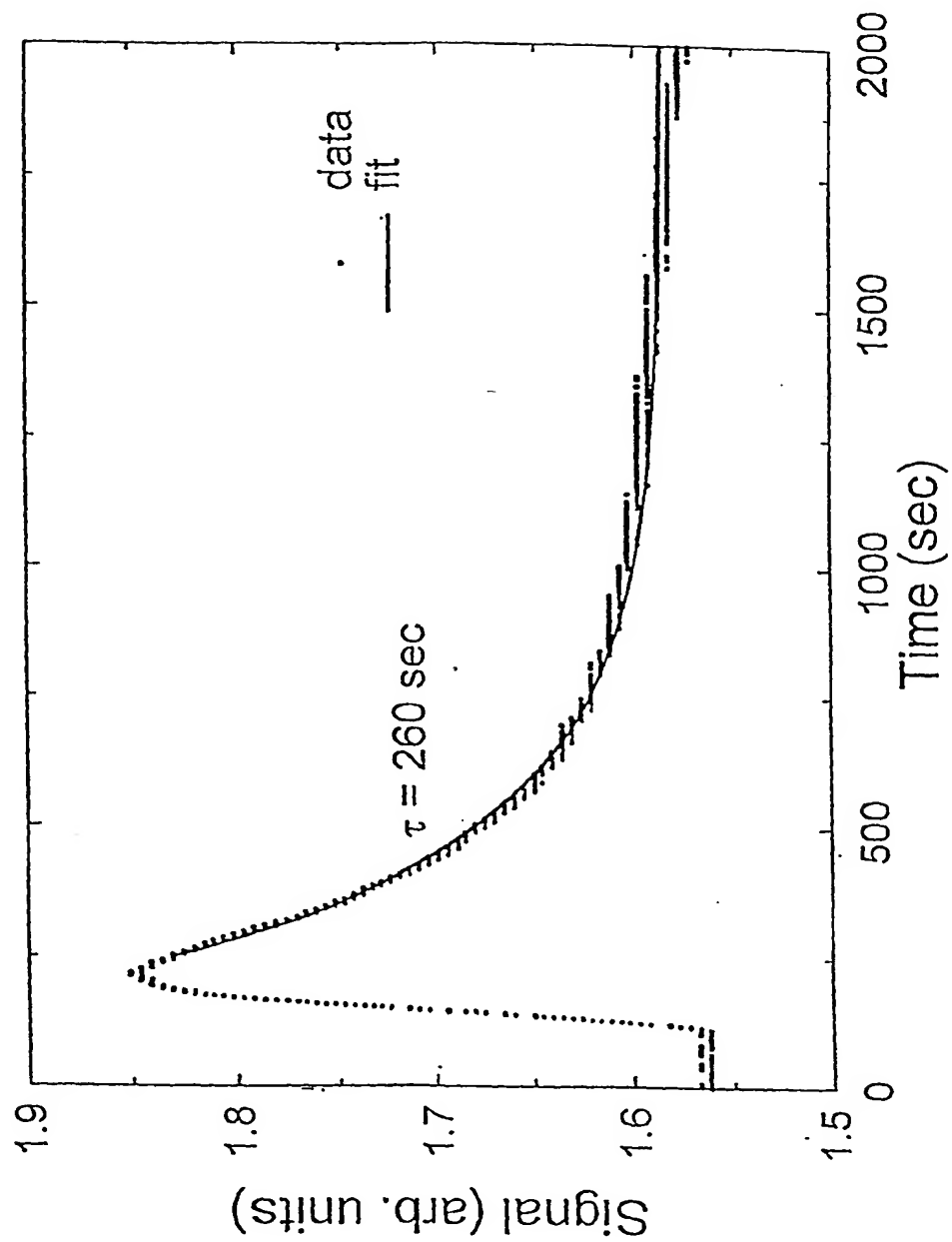


Figure 12

Blood clearance profile of indole disulfonate

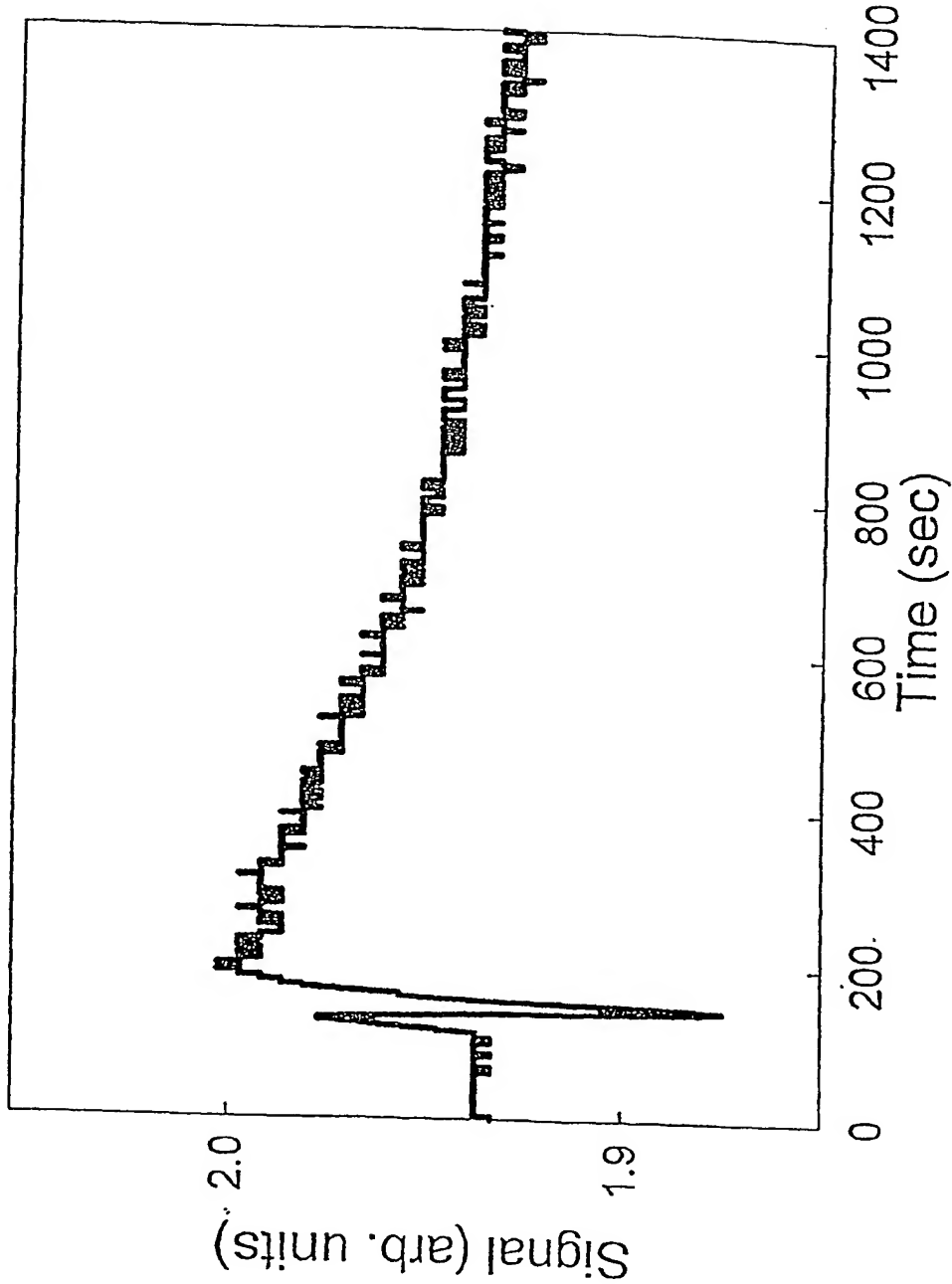


Figure 13

# Blood clearance profile of cyaninetetrasulfonates

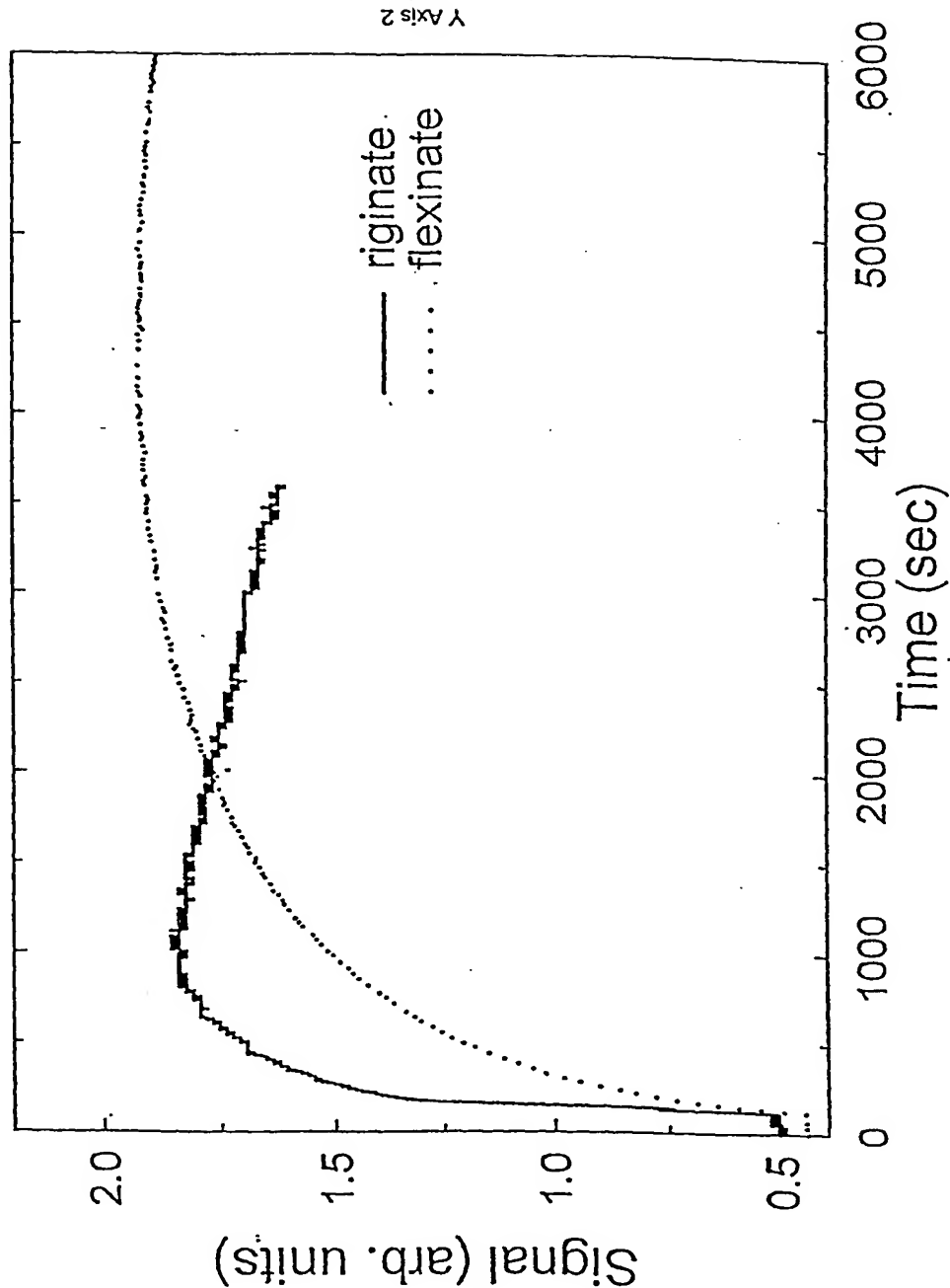


Figure 14

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/31718

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A61K 49/00; C07D 209/04

US CL : 252/408.1; 548/469

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
U.S. : 252/408.1; 548/469Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched  
NONEElectronic data base consulted during the international search (name of data base and, where practicable, search terms used)  
STN: structure search

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X --- A	US 5,453,505 A (LEE et al) 26 September 1995 (26.09.95), see abstract and Table 1, NHCL dye.	1 ----- 2-14
X --- A	Registry No. 259261-66-0, see structure.	1 ----- 2-14

☐ Further documents are listed in the continuation of Box C.☐ See patent family annex.

Special categories of cited documents	
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"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

05 March 2002 (05.03.2002)

Date of mailing of the international search report

04 APR 2002

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Facsimile No. (703)305-3230

Authorized officer

Leigh McKane

Telephone No. 703-308-0661

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